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# (54) Title: DEVELOPMENT OF NOVEL ANTI-MICROBIAL AGENTS BASED ON BACTERIOPHAGE GENOMICS

#### (57) Abstract

A method for identifying suitable targets for antibacterial agents based on identifying targets of bacteriophage-encoded proteins is described. Also described are compositions useful in the identification methods and in inhibiting bacterial growth, and methods for preparing and using such compositions.

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1

# **DESCRIPTION**

# Development of Novel Anti-Microbial Agents Based on Bacteriophage Genomics

# **BACKGROUND OF THE INVENTION**

The present invention relates to the field of antibacterial agents and the treatment of infections of animals or other complex organisms by bacteria.

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The frequency and spectrum of antibiotic-resistant infections have, in recent years, increased in both the hospital and community. Certain infections have become essentially untreatable and are growing to epidemic proportions in the developing world as well as in institutional settings in the developed world. The staggering spread of antibiotic resistance in pathogenic bacteria has been attributed to microbial genetic characteristics, widespread use of antibiotic drugs, and changes in society that enhance the transmission of drug-resistant organisms. This spread of drug resistant microbes is leading to ever increasing morbidity, mortality and health-care costs.

Ironically, it is the very success of antibiotics, resulting in their widespread use, that has contributed the most to rising numbers of drug resistant bacterial strains. The longer a bacterial strain is exposed to a drug, the more likely it is to acquire resistance. Today, a total of 160 antibiotics, all based on a few basic chemical structures and targeting a small number of metabolic pathways, have found their way to market. Over-prescription of these drugs, as well as the failure of patients to comply with the complete antibiotic regimen, has lead to the rapid emergence of antibiotic resistant strains. Such misuse of prescriptions, careless use of antibiotics in virtually all commercial production of beef and fowl, and changing societal conditions, such as the growth of day-care centers, increased long-term care in hospitals, and increased mobility of the population, has provided an environment where drug-resistant microbes can emerge and spread. Thus, virtually all common infectious bacteria are becoming, or have already become, resistant to one or more groups of antibiotics. Such resistance now reaches all classes of antibiotics currently in use, including: β-lactams, fluoroquinolones, aminoglycosides, macrolide peptides, chloramphenicol, tetracyclines, rifampicin, folate inhibitors, glycopeptides, and mupirocin.

Over the last 45 years bacteria have adapted genetically to avoid the destruction/alteration of the essential pathways that these chemotherapeutic agents

2

target. Antibiotic resistant bacterial strains are now emerging at a higher rate than the rate at which new antibiotics are being developed. The consequence of this dilemma has been a dramatic increase in the cost of treating infections what would otherwise easily succumb to routine antibiotic therapy. Furthermore, and perhaps most importantly, the emergence of multiple drug resistant pathogenic bacteria has led to a significant increase in morbidity and mortality, particularly in institutional settings.

Most major pharmaceutical companies have on-going drug discovery programs for novel anti-microbials. These are based on screens for small molecule inhibitors (natural products, bacterial culture media, libraries of small molecules, combinatorial chemistry) of crucial metabolic pathways of the micro-organism of interest (e.g., bacteria, fungi, parasites, worms). The screening process is largely for cytotoxic compounds and in most cases is not based on a known mechanism of action of the compounds. Pharmaceutical companies have large programs in this area. Classical drug screening programs are being exhausted and many of these pharmaceutical companies are looking towards rational drug design programs.

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Several small to mid-size biotechnology companies as well as large pharmaceutical companies have developed systematic high-throughput sequencing programs to decipher the genetic code of specific micro-organisms of interest. The goal is to identify, through sequencing, unique biochemical pathways or intermediates that are unique to the microorganism. Knowledge of this may, in turn, form the rationale for a drug discovery program based on the mechanism of action of the identified enzymes/proteins. Genome Therapeutics Corp., The Institute for Genome Research, Human Genome Sciences Inc., and other companies have such sequencing programs in place. However, one of the most critical steps in this approach is the ascertainment that the identified proteins and biochemical pathways are 1) non-redundant and essential for bacterial survival, and 2) constitute suitable and accessible targets for drug discovery.

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# SUMMARY OF THE INVENTION

While animals such as humans are, on occasion, infected by pathogenic bacteria, bacteria also have natural enemies. A number of host-specific viruses, known as bacteriophages or phages, infect and kill bacteria in the natural environment. Such bacteriophages generally have small compact genomes and bacteria are their exclusive hosts. Many known bacteria are host to a large number of bacteriophages that have been described in the literature. During the 1940's - 1960's, phage biology was an area of active research. As a testimony to this, the study of phages which infect and inhibit the enteric bacterium *Escherichia coli* (*E. coli*) contributed much to the early understanding of molecular biology and virology.

As is generally understood, bacteriophage (or phages) are viruses that infect and kill bacteria. They are natural enemies of bacteria and, over the course of evolution, have developed proteins (products of DNA sequences) which enable them to infect a host bacteria, replicate their genetic material, usurp host metabolism, and ultimately kill their host. The scientific literature well documents the fact that many known bacteria have a large number of such bacteriophages (Ackermann and DuBow, 1987) that can infect and kill them (for example, see the ATCC bacteriophage collection at http://www.atcc.org).

This invention utilizes the observation that bacteriophages successfully infect and inhibit or kill host bacteria, targeting a variety of normal host metabolic and physiological traits, some of which are shared by all bacteria, pathogenic and nonpathogenic alike. The term "pathogenic" as used herein denotes a contribution to or implication in disease or a morbid state of an infected organism. The invention thus involves identifying and elucidating the molecular mechanisms by which phages interfere with host bacterial metabolism, an objective being to provide novel targets for drug design. Whether the phage blocks bacterial RNA transcription or translation, or attacks other important metabolic pathways, such as cell wall assembly or membrane integrity, the basic blueprint for a phage's bacteria-inhibiting ability is encoded in its genome and can be unlocked using bioinformatics, functional genomics, and proteomics. By these means, the invention utilizes sequence information from the genomics of bacteriophage to identify novel antimicrobials that can be further used to actively and/or prophylactically treat bacterial infection.

Two important components of the invention thus are: i) the identification of bacteria-inhibiting phage open reading frames ("ORF"s) and corresponding products that can be used to develop antibiotics based on amino acid sequence and secondary structural characteristics of the ORF products, and ii) the use of bacteriophages to map

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out essential bacterial target genes and homologs, which can in turn lead to the development of suitable anti-microbial agents. These two avenues represent new and general methods for developing novel antimicrobials.

The invention thus concerns the identification of bacteriophage ORFs that supply bacteria-inhibiting functions. In this regard, use of the terms "inhibit", "inhibition", "inhibitory", and "inhibitor" all refer to a function of reducing a biological activity or function. Such reduction in activity or function can, for example, be in connection with a cellular component, e.g., an enzyme, or in connection with a cellular process, e.g., synthesis of a particular protein, or in connection with an overall process of a cell, e.g., cell growth. In reference to bacterial cell growth, for example, an inhibitory effect (i.e., a bacteria-inhibiting effect) may be bacteriocidal (killing of bacterial cells) or bacteriostatic (i.e., stopping or at least slowing bacterial cell growth). The latter slows or prevents cell growth such that fewer cells of the strain are produced relative to uninhibited cells over a given period of time. From a molecular standpoint, such inhibition may equate with a reduction in the level of, or elimination of, the transcription and/or translation of a specific bacterial target(s), or reduction or elimination of activity of a particular target biomolecule.

It is particularly advantageous to evaluate a plurality of different phage ORFs for inhibitory activity that may be from one, but is preferably from a plurality of different phage. For example, evaluating ORFs from a number of different phage of the same bacterial host provides at least two advantages. One is that the multiple phages will provide identification of a variety of different targets. Second, it is likely that multiple phage will utilize the same cellular target

As used herein, the terms "bacteriophage" and "phage" are used interchangeably to refer to a virus which can infect a bacterial strain or a number of different bacterial strains.

In the context of this invention, the term "bacteriophage ORF" or "phage ORF" or similar term refers to a nucleotide sequence in or from a bacteriophage. In connection with a particular ORF, the terms refer an open reading frame which has at least 95% sequence identity, preferably at least 97% sequence identity, more preferably at least 98% sequence identity with an ORF from the particular phage identified herein (e.g., with an ORF as identified herein) or to a nucleic acid sequence which has the specified sequence identify percentage with such an ORF sequence.

A first aspect of the invention thus provides a method for identifying a bacteriophage nucleic acid coding region encoding a product active on an essential bacterial target by identifying a nucleic acid sequence encoding a gene product which

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provides a bacteria-inhibiting function when the bacteriophage infects a host bacterium, preferably one that is an animal or plant pathogen, more preferably a bird or mammalian pathogen, and most preferably a human pathogen. The bacteriophage is an uncharacterized bacteriophage. Thus, the method excludes, for example, phage  $\lambda$ ,  $\phi$ x174, m13 and other *E.coli*-specific bacteriophage that have been studied with respect to gene number and/or function. It also excludes, for example, the nucleic acid coding regions described in Tables 12-14, and in preferred embodiments, excludes the phage in which those regions are naturally located.

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In connection with bacteriophage, the term "uncharacterized" means that a certain bacteriophage's genome has not yet been fully identified such that the genes having function involved in inhibiting host cells have not been identified. In particular, phage for which the description of genomic or protein sequence was first provided herein are uncharacterized. Phage sequences for which host bacteriainhibiting functions have been identified prior to the filing of the present application (or alternatively prior to the present invention) are specifically excluded from the aspects involving utilization of sequences from uncharacterized bacteriophage, except that aspects may involve a plurality of phage where one or more of those phage are uncharacterized and one or more others have been characterized to some extent. A number of different bacteria-inhibiting phage ORFs are indicated in Tables 11-14. The phage ORFs or sequences identified therein are not within the term "uncharacterized; alternatively, in preferred embodiments the phage containing those ORFs are excluded from this term. Further, any additional phage ORFs (or alternatively the phage which contain those ORFs) which have previously been described in the art as bacteria-inhibiting ORFs are expressly excluded; those ORFs or phage are known to those skilled in the art and the exclusion can be made express by specifically naming such ORFs or phage as needed (likewise for uncharacterized targets as described below). For the sake of brevity, such a listing is not expressly

Stating that an agent or compound is "active on" a particular cellular target, such as the product of a particular gene, means that the target is an important part of a cellular pathway which includes that target and that the agent acts on that pathway. Thus, in some cases the agent may act on a component upstream or downstream of the stated target, including on a regulator of that pathway or a component of that pathway.

presented, as such information is readily available to those skilled in the art.

By "essential", in connection with a gene or gene product, is meant that the host cannot survive without, or is significantly growth compromised, in the absence depletion, or alteration of functional product. An "essential gene" is thus one that encodes a product that is beneficial, or preferably necessary, for cellular growth in

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vitro in a medium appropriate for growth of a strain having a wild-type allele corresponding to the particular gene in question. Therefore, if an essential gene is inactivated or inhibited, that cell will grow significantly more slowly, preferably less than 20%, more preferably less than 10%, most preferably less than 5% of the growth rate of the uninhibited wild-type, or not at all, in the growth medium. Preferably, in the absence of activity provided by a product of the gene, the cell will not grow at all or will be non-viable, at least under culture conditions similar to the *in vivo* conditions normally encountered by the bacterial cell during an infection. For example, absence of the biological activity of certain enzymes involved in bacterial cell wall synthesis can result in the lysis of cells under normal osmotic conditions, even though protoplasts can be maintained under controlled osmotic conditions. In the context of the invention, essential genes are generally the preferred targets of antimicrobial agents. Essential genes can encode target molecules directly or can encode a product involved in the production, modification, or maintenance of a target molecule.

A "target" refers to a biomolecule that can be acted on by an exogenous agent, thereby modulating, preferably inhibiting, growth or viability of a cell. In most cases such a target will be a nucleic acid sequence or molecule, or a polypeptide or protein. However, other types of biomolecules can also be targets, e.g., membrane lipids and cell wall structural components.

The term "bacterium" refers to a single bacterial strain, and includes a single cell, and a plurality or population of cells of that strain unless clearly indicated to the contrary. In reference to bacteria or bacteriophage, the term "strain" refers to bacteria or phage having a particular genetic content. The genetic content includes genomic content as well as recombinant vectors. Thus, for example, two otherwise identical bacterial cells would represent different strains if each contained a vector, e.g., a plasmid, with different phage ORF inserts.

In preferred embodiments, the phage is *Staphylococcus aureus* phage 77, 3A, 96, or 44 AHJD, *Enterococcus* sp. phage 182, or *Streptococcus pneumoniae* phage Dp-1.

In preferred embodiments, the phage is selected from. Preferred embodiments involve expressing at least one recombinant phage ORF(s) in a bacterial host followed by inhibition analysis of that host. Inhibition following expression of the phage ORF is indicative that the product of the ORF is active on an essential bacterial target. Such evaluation can be carried out in a variety of different formats, such as on a support matrix such as a solidified medium in a petri dish, or in liquid culture.

7

Preferably a plurality of phage ORFs are expressed in at least one bacterium. The plurality of phage ORFs can be from one or a plurality of phage. With respect to a single phage or at least one phage in a plurality of phages, the plurality of expressed ORFs preferably represents at least 10%, more preferably at least 20%, 40%, or 60%, still more preferably at least 80% or 90%, and most preferably at least 95% of the ORFs in the phage genome. Preferably, for a plurality of phage, the plurality of expressed ORFs preferably represents at least 10%, more preferably at least 20%, 40%, or 60%, still more preferably at least 80% or 90%, and most preferably at least 95% of the ORFs in the phage genome of each phage. The plurality of phage ORFs can be expressed in a single bacterium, or in a plurality of bacteria where one ORF is expressed in each bacterium, or in a plurality of bacteria where a plurality of ORFs are expressed in at least one or in all of the plurality of bacteria, or combinations of these.

In embodiments of the above aspect (as well as in other aspects herein) in which a plurality of phage are utilized, a plurality of phage have the same bacterial host species; have different bacterial host species; or both. The plurality of phage includes at least two different phage, preferably at least 3,4,5,6,8,10,15,20, or more different phage. Indeed, more preferably, the plurality of phage will include 50, 75, 100, or more phage. As described herein, the larger number of phage is useful to provide additional target and target evaluation information useful in developing antibacterial agents, for example, by providing identification of a larger range of bacterial targets, and/or providing further indication of the suitability of a particular target (for example, utilization of a target by a number of different unrelated phage can suggest that the target is particularly stable and accessible and effective) and/or can indicate alternate sites on a target which interact with different inhibitors.

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Further embodiments involve confirmation of the inhibitor function of the phage ORF, such as by utilizing or incorporating a control(s) designed to confirm the inhibitory nature of the ORF(s) being evaluated. The control can, for example, be provided by expression of an inactive or partially inactive form of the ORF or ORF product, and/or by the absence of expression of the ORF or ORF product in the same or a closely comparable bacterial strain as that used for expression of the test ORF. The reduced level of activity or the absence of active ORF product in the control will thus not provide the inhibition provided by a corresponding inhibitory ORF, or will provide a distinguishably lower level of inhibition. An inactivated or partially inactivated control has a mutation(s), e.g., in the coding region or in flanking regulatory elements, that reduce(s) or eliminate(s) the normal function of the ORF. Thus, the inhibition of a bacterium following expression of a phage ORF is determined by comparison with the effects of expression of an inactivated ORF or the

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response of the bacteria in the absence of expression in the same or similar type bacterium. Such determination of inhibition of the bacterium following expression of the ORF is indicative of a bacteria-inhibiting function. These manipulations are routinely understood and accomplished by those of skill in the art using standard techniques. In embodiments utilizing absence of expression of the ORF, the bacteria can, for example, contain an empty vector or a vector which allows expression of an unrelated sequence which is preferably non-inhibitory. Alternatively, the bacteria may have no vector at all. Combinations of such controls or other controls may also be utilized as recognized by those skilled in the art.

In embodiments involving expression of a phage ORF in a bacterial strain, in preferred embodiments that expression is inducible.

By "inducible" is meant that expression is absent or occurs at a low level until the occurrence of an appropriate environmental stimulus provides otherwise. For the present invention such induction is preferably controlled by an artificial environmental change, such as by contacting a bacterial strain population with an inducing compound (i.e., an inducer). However, induction could also occur, for example, in response to build-up of a compound produced by the bacteria in the bacterial culture, e.g., in the medium. As uncontrolled or constitutive expression of inhibitory ORFs can severely compromise bacteria to the point of eradication, such expression is therefore undesirable in many cases because it would prevent effective evaluation of the strain and inhibitor being studied. For example, such uncontrolled expression could prevent any growth of the strain following insertion of a recombinant ORF, thus preventing determination of effective transfection or transformation. A controlled or inducible expression is therefore advantageous and is generally provided through the provision of suitable regulatory elements, e.g., promoter/operator sequences that can be conveniently transcriptionally linked to a coding sequence to be evaluated. In most cases, the vector will also contain sequences suitable for efficient replication of the vector in the same or different host cells and/or sequences allowing selection of cells containing the vector, i.e., "selectable markers." Further, preferred vectors include convenient primer sequences flanking the cloning region from which PCR and/or sequencing may be performed.

As knowledge of the nucleotide sequence of phage ORFs is useful, e.g., for assisting in the identification of phage proteins active against essential bacterial host targets, preferred embodiments involve the sequencing of at least a portion of the phage genome in combination with the above methods. This can be done either before or after or independent of expression and inhibition of the ORF in the bacteria, and provides information on the nature and characteristics of the ORF. Such a portion is

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preferably at least 10%, 20%, 40%, 80%, 90%; or 100% of the phage genome. For embodiments in which a plurality of phage are utilized, preferably each phage is sequenced to an extent as just specified.

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Such sequencing is preferably accompanied by computer sequence analysis to define and evaluate ORF(s), ORF products, structural motifs or functional properties of ORF products, and/or their genetic control elements. Thus, certain embodiments incorporate computer sequence analyses or nucleic acid and/or amino acid sequences. Further, existing data banks can provide phage sequence and product information which can be utilized for analysis and identification of ORFs in the sequence.

Computer analysis may further employ known homologous sequences from other species that suggest or indicate conserved underlying biochemical function(s) for the inhibitory or potentially inhibitory ORF sequence(s) being evaluated. This can include the sequences of signature motifs of identified classes of inhibitors.

In the context of the phage nucleic acid sequences, e.g., gene sequences, of this invention, the terms "homolog" and "homologous" denote nucleotide sequences from different bacteria or phage strains or species or from other types of organisms that have significantly related nucleotide sequences, and consequently significantly related encoded gene products, preferably having related function. Homologous gene sequences or coding sequences have at least 70% sequence identity (as defined by the maximal base match in a computer-generated alignment of two or more nucleic acid sequences) over at least one sequence window of 48 nucleotides, more preferably at least 80 or 85%, still more preferably at least 90%, and most preferably at least 95%. The polypeptide products of homologous genes have at least 35% amino acid sequence identity over at least one sequence window of 18 amino acid residues, more preferably at least 40%, still more preferably at least 50% or 60%, and most preferably at least 70%, 80%, or 90%. Preferably, the homologous gene product is also a functional homolog, meaning that the homolog will functionally complement one or more biological activities of the product being compared. For nucleotide or amino acid sequence comparisons where a homology is defined by a % sequence identity, the percentage is determined using BLAST programs ( with default parameters (Altschul et al., 1997, "Gapped BLAST and PSI-BLAST: a new generation of protein database search programs, Nucleic Acid Res. 25:3389-3402). Any of a variety of algorithms known in the art which provide comparable results can also be used, preferably using default parameters. Performance characteristics for three different algorithms in homology searching is described in Salamov et al., 1999, "Combining sensitive database searches with multiple intermediates to detect distant

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homologues." *Protein Eng.* 12:95-100. Another exemplary program package is the  $GCG^{TM}$  package from the University of Wisconsin.

Homologs may also or in addition be characterized by the ability of two complementary nucleic acid strands to hybridize to each other under appropriately stringent conditions. Hybridizations are typically and preferably conducted with probe-length nucleic acid molecules, preferably 20-100 nucleotides in length. Those skilled in the art understand how to estimate and adjust the stringency of hybridization conditions such that sequences having at least a desired level of complementarity will stably hybridize, while those having lower complementarity will not. For examples of hybridization conditions and parameters, see, e.g., Maniatis, T. et al. (1989)

Molecular Cloning: A Laboratory Manual, Cold Spring Harbor University Press, Cold Spring, N.Y.; Ausubel, F.M. et al. (1994) Current Protocols in Molecular Biology.

John Wiley & Sons, Secaucus, N.J. Homologs and homologous gene sequences may thus be identified using any nucleic acid sequence of interest, including the phage ORFs and bacterial target genes of the present invention.

A typical hybridization, for example, utilizes, besides the labeled probe of interest, a salt solution such as 6xSSC (NaCl and Sodium Citrate base) to stabilize nucleic acid strand interaction, a mild detergent such as 0.5% SDS, together with other typical additives such as Denhardt's solution and salmon sperm DNA. The solution is added to the immobilized sequence to be probed and incubated at suitable temperatures to preferably permit specific binding while minimizing nonspecific binding. The temperature of the incubations and ensuing washes is critical to the success and clarity of the hybridization. Stringent conditions employ relatively higher temperatures, lower salt concentrations, and/or more detergent than do non-stringent conditions. Hybridization temperatures also depend on the length, complementarity level, and nature (ie, "GC content") of the sequences to be tested. Typical stringent hybridizations and washes are conducted at temperatures of at least 40°C, while lower stringency hybridizations and washes are typically conducted at 37°C down to room temperature (~25°C). One of skill in the art is aware that these conditions may vary according to the parameters indicated above, and that certain additives such as formamide and dextran sulphate may also be added to affect the conditions.

By "stringent hybridization conditions" is meant hybridization conditions at least as stringent as the following: hybridization in 50% formamide, 5X SSC, 50 mM NaH,PO., pH 6.8, 0.5% SDS, 0.1 mg/mL sonicated salmon sperm DNA, and 5X Denhart's solution at 42°C overnight; washing with 2X SSC, 0.1% SDS at 45°G; and washing with 0.2X SSC, 0.1% SDS at 45°C.

11

In sequence comparison analyses, an ORF, or motif, or set of motifs in a bacteriophage sequence can be compared to known inhibitor sequences, e.g., homologous sequences encoding homologous inhibitors of bacterial function. Likewise, the analysis can include comparison with the structure of essential bacterial gene products, as structural similarities can be indicative of similar or replacement biological function. Such analysis can include the identification of a signature, or characteristic motif(s) of an inhibitor or inhibitor class.

Also, the identification of structural motifs in an encoded product, based on nucleotide or amino acid sequence analysis, can be used to infer a biochemical function for the product. A database containing identified structural motifs in a large number of sequences is available for identification of motifs in phage sequences. The database is PROSITE, which is available at www.expasy.ch/cgi~bin/scanprosite. The identification of motifs can, for example, include the identification of signature motifs for a class or classes of inhibitory proteins. Other such databases may also be used.

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In aspects and preferred embodiments described herein, in which a bacterium or host bacterium is specified, the bacterium or host bacterium is preferably selected from a pathogenic bacterial species, for example, one selected from Table 1. Preferably, an animal or plant pathogen is used. For animals, preferably the bacterium is a bird or mammalian pathogen, still more preferably a human pathogen.

In aspects and preferred embodiments involving a bacteriophage or sequences from a bacteriophage, one or more bacteriophage are preferably selected from those listed in Table 1. Those exemplary bacteriophage are readily obtained from the indicated sources.

In some cases, it is advantageous to utilize phage with non-pathogenic host bacteria. The genome, structural motif, ORF, homolog, and other analyses described herein can be performed on such phage and bacteria. Such analysis provides useful information and compositions. The results of such analyses can also be utilized in aspects of the present invention to identify homologous ORFs, especially inhibitor ORFs in phage with pathogenic bacterial hosts. Similarly, identification of a target in a non-pathogenic host can be used to identify homologous sequences and targets in pathogenic bacteria, especially in genetically closely related bacteria. Those skilled in the art are familiar with bacterial genetic relationships and with how to determine relatedness based on levels of genomic identity or other measures of nucleotide sequence and/or amino acid sequence similarity, and/or other physical and culture characteristics such as morphology, nutritional requirements, or minimal media to support growth.

Also in preferred embodiments, an embodiments of this aspect is combined with an embodiment of the following aspect.

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A related aspect of the invention provides methods for identifying a target for antibacterial agents by identifying the bacterial target(s) of at least one uncharacterized or untargeted inhibitor protein or RNA from a bacteriophage. Such identification allows the development of antibacterial agents active on such targets. Preferred embodiments for identifying such targets involve the identification of binding of target and phage ORF products to one another. The phage ORF products may be subportions of a larger ORF product that also binds the host target. In preferred embodiments, the phage protein or RNA is from an uncharacterized bacteriophage in Table 1. This aspect preferably includes the identification of a plurality of such targets in one or a plurality of different bacteria, preferably in one or a plurality of bacteria listed in Table 1.

In preferred embodiments of this aspect and other aspects of this invention involving particular phage ORFs or phage sequences, the ORF is *Staphylococcus aureus* phage 77 ORF 17, 19, 43, 102, 104, or 182 as identified in U.S. application 09/407,804, *S. aureus* phage 44AHJD ORF 1, 9, or 12, *Streptococcus pneumoniae* phage Dp-1 ORF 001, 002, 004, 008, 010, 013, 016, 021, 029, 030, 038, or 041, or *Enterococcus* sp. phage 182 ORF 002, 008, or 014.

As indicated for the above aspect, preferably the method involves the use of a plurality of different phage, and thus a plurality of different phage inhibitors and/or inhibitor ORFs.

In addition to uncharacteized phage ORF products, it is also useful to identify the targets of phage ORF products which are known to be inhibitors of host bacteria, but where the target has not been identified. Thus, such inhibitors can likewise be utilized as "untargeted" inhibitor phage ORFs and ORF products, e.g., proteins or RNAs.

In the context of inhibitor proteins or RNAs from a phage, the term "uncharacterized" means that a bacteria-inhibiting function for the protein has not previously been identified. Preferably, but not necessarily, the sequence of the protein or the corresponding coding region or ORF was not described in the art before the filing of the present application for patent (or alternatively prior to the present invention). Thus, this term specifically excludes any bacteria-inhibiting phage protein and its associated bacterial target which has been identified as inhibitory before the present invention or alternatively before the filing of the present application, for. example those identified in Tables 12-14 or otherwise identified herein. For example, from *E. coli*, phage T7 genes 0.7 and 2.0 target the host RNA polymerase, phage T4

13

gp55/gp33 alter the specificity of host RNA polymerase. The T4 regB gene product also targets the host translation apparatus. As with the uncharacterized bacteriophage ORFs or bacteriophage above, for such identified proteins, the sequences encoding those proteins are excluded from the uncharacterized inhibitor proteins.

The term "fragment" refers to a portion of a larger molecule or assembly. For proteins, the term "fragment" refers to a molecule which includes at least 5 contiguous amino acids from the reference polypeptide or protein, preferably at least 8, 10, 12, 15, 20, 30, 50 or more contiguous amino acids. In connection with oligo- or polynucleotides, the term "fragment" refers to a molecule which includes at least 15 contiguous nucleotides from a reference polynucleotide, preferably at least 24, 30, 36, 45, 60, 90, 150, or more contiguous nucleotides.

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Preferred embodiments involve identification of binding that include methods for distinguishing bound molecules, for example, affinity chromatography, immunoprecipitation, crosslinking, and/or genetic screen methods that permit protein:protein interactions to be monitored. One of skill in the art is familiar with these techniques and common materials utilized (see, e.g., Coligan, J. et al. (eds.) (1995) Current Protocols in Protein Science, John Wiley & Sons, Secaucus, N.J.).

Genetic screening for the identification of protein:protein interactions typically involves the co-introduction of both a chimeric bait nucleic acid sequence (here, the phage ORF to be tested) and a chimeric target nucleic acid sequence that, when co-expressed and having affinity for one another in a host cell, stimulate reporter gene expression to indicate the relationship. A "positive" can thus suggest a potential inhibitory effect in bacteria. This is discussed in further detail in the Detailed Description section below. In this way, new bacterial targets can be identified that are inhibited by specific phage ORF products or derivatives, fragments, mimetics, or other molecules.

Other embodiments involve the identification and/or utilization of mutant targets by virtue of their host's relatively unresponsive nature in the presence of expression of ORFs previously identified as inhibitory to the non-mutant or wild-type strain. Such mutants have the effect of protecting the host from an inhibition that would otherwise occur and indirectly allow identification of the precise responsible target for follow-up studies and anti-microbial development. In certain embodiments, rescue from inhibition occurs under conditions in which a bacterial target or mutant target is highly expressed. This is performed, for example, through coupling of the sequence with regulatory element promoters, e.g., as known in the art, which regulate expression at levels higher than wild-type, e.g., at a level sufficiently higher that the

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inhibitor can be competitively bound to the highly expressed target such that the bacterium is detectably less inhibited.

Identification of the bacterial target can involve identification of a phage-specific site of action. This can involve a newly identified target, or a target where the phage site of action differs from the site of action of a previously known antibacterial agent or inhibitor. For example, phage T7 genes 0.7 and 2.0 target the host RNA polymerase, which is also the cellular target for the antibacterial agent, rifampin. To the extent that a phage product is found to act at a different site than previously described inhibitors, aspects of the present invention can utilize those new, phage-specific sites for identification and use of new agents. The site of action can be identified by techniques well-known to those skilled in the art, for example, by mutational analysis, binding competition analysis, and/or other appropriate techniques.

Once a bacterial host target protein or nucleic acid or mutant target sequence has been identified and/or isolated, it too can be conveniently sequenced, sequence analyzed (e.g., by computer), and the underlying gene(s), and corresponding translated product(s) further characterized. Preferred embodiments include such analysis and identification. Preferably such a target has not previously been identified as an appropriate target for antibacterial action.

Certain embodiments include the identification of at least one inhibitory phage ORF or ORF product, e.g., as described for the above aspect, and thus are a combination of the two aspects.

Additionally, the invention provides methods for identifying targets for antibacterial agents by identifying homologs of a bacterial target e.g., S. aureus, Enterococcus faecalis or other Enterococci, and Streptococcus pneumoniae of a bacteriophage inhibitory ORF product. Such homologs may be utilized in the various aspects and embodiments described herein as described for the host Enterococcus sp. for bacteriophage 182.

Other aspects of the invention provide isolated, purified, or enriched specific phage nucleic acid and amino acid sequences, subsequences, and homologs thereof for phage selected from uncharacterized phage listed in Table 1, preferably from bacteriophage 77, 3A, 96, 44AHJD (Staphylococcus aureus host bacterium), Dp-1 (Streptococcus pneumoniae host), or 182 (Enterococcus host) or other phage listed in Table 1 for those bacteria. For example, such sequences do not include sequences identified in any of Tables 11-14. Nucleotide sequences of this aspect are at least 15 nucleotides in length, preferably at least 18, 21, 24, or 27 nucleotides in length, more preferably at least 30, 50, or 90 nucleotides in length. In certain embodiments, longer

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nucleic acids are preferred, for example those of at least 120, 150, 200, 300, 600, 900 or more nucleotides. Such sequences can, for example, be amplification oligonucleotides (e.g., PCR primers), oligonucleotide probes, sequences encoding a portion or all of a phage-encoded protein, or a fragment or all of a phage-encoded protein. In preferred embodiments, the nucleic acid sequence contains a sequence which is within a length range with a lower length as specified above, and an upper length limit which is no more than 50, 60, 70, 80, or 90% of the length of the corresponding full-length ORF. The upper length limit can also be expressed in terms of the number of base pairs of the ORF (coding region). In preferred embodiments, the nucleic acid sequence is from Staphylococcus aureus phage 77 ORF 17, 19, 43, 102, 104, or 182 as identified in U.S. application 09/407,804, S. aureus phage 44 AHJD ORF 1, 9, or 12, Streptococcus pneumoniae phage Dp-1 ORF 001, 002, 004, 008, 010, 013, 016, 021, 029, 030, 038, or 041, or Enterococcus sp. phage 182 ORF 002, 008, or 014.

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As it is recognized that alternate codons will encode the same amino acid for most amino acids due to the degeneracy of the genetic code, the sequences of this aspect includes nucleic acid sequences utilizing such alternate codon usage for one or more codons of a coding sequence. For example, all four nucleic acid sequences GCT, GCC, GCA, and GCG encode the amino acid, alanine. Therefore, if for an amino acid there exists an average of three codons, a polypeptide of 100 amino acids in length will, on average, be encoded by 3100, or 5 x 1047, nucleic acid sequences. Thus, a nucleic acid sequence can be modified (e.g., a nucleic acid sequence from a phage as specified above) to form a second nucleic acid sequence encoding the same polypeptide as encoded by the first nucleic acid sequence using routine procedures and without undue experimentation. Thus, all possible nucleic acid sequences that encode the specified amino acid sequences are also fully described herein, as if all were written out in full, taking into account the codon usage, especially that preferred in the host bacterium. The alternate codon descriptions are available in common texbooks, for example, Stryer, BIOCHEMISTRY 3rd ed., and Lehninger, BIOCHEMISTRY 3rd ed., along wth many others. Codon preference tables for various types of organisms are available in the literature. Sequences with alternate codons at one or more sites can also be utilized in the computer-related aspects and embodiments herein. Because of the number of sequence variations involving alternate codon usage, for the sake of brevity, individual sequences are not separately listed herein. Instead the alternate sequences are described by reference to the natural sequence with replacement of one or more (up to all e.g., up to 3, 5, 10, 15, 20, 30, 40, 50, or more) of the degenerate codons with alternate codons from the alternate codon

16

table (Table 6), or a modified table applicable to a particular organism that has differing codon usage, preferably with selection according to preferred codon usage for the normal host organism or a host organism in which a sequence is intended to be expressed. Those skilled in the art also understand how to alter the alternate codons to be used for expression in organisms where certain codons code differently than shown in the "universal" codon table.

For amino acid sequences or polypeptides, sequences contain at least 5 peptide-linked amino acid residues, and preferably at least 6, 7, 10, 15, 20, 30, or 40, amino acids having identical amino acid sequence as the same number of contiguous amino acid residues in a particular phage ORF product. In some cases longer sequences may be preferred, for example, those of at least 50, 60, 70, 80, or 100 amino acids in length. In preferred embodiments, the amino acid sequence contains a sequence which is within a length range with a lower length as specified above, and an upper length limit which is no more than 50, 60, 70, 80, or 90% of the length of the corresponding full-length ORF product. The upper length limit can also be expressed in terms of the number of amino acid residues of the ORF product. In preferred embodiments, the amino acid sequence or polypeptide has bacteria-inhibiting function when expressed or otherwise present in a bacterial cell which is a host for the bacteriophage from which the sequence was derived.

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By "isolated" in reference to a nucleic acid is meant that a naturally occurring sequence has been removed from its normal cellular (e.g., chromosomal) environment or is synthesized in a non-natural environment (e.g., artificially synthesized). Thus, the sequence may be in a cell-free solution or placed in a different cellular environment. The term does not imply that the sequence is the only nucleotide chain present, but that it is essentially free (about 90-95% pure at least) of non-nucleotide material naturally associated with it, and thus is distinguished from isolated chromosomes.

The term "enriched" means that the specific DNA or RNA sequence constitutes a significantly higher fraction (2-5 fold) of the total DNA or RNA present in the cells or solution of interest than in normal or diseased cells or in cells from which the sequence was originally taken. This could be caused by a person by preferential reduction in the amount of other DNA or RNA present, or by a preferential increase in the amount of the specific DNA or RNA sequence, or by a combination of the two. However, it should be noted that enriched does not imply that there are no other DNA or RNA sequences present, just that the relative amount of the sequence of interest has been significantly increased.

17

The term "significant" is used to indicate that the level of increase is useful to the person making such an increase and an increase relative to other nucleic acids of about at least 2-fold, more preferably at least 5- to 10-fold or even more. The term also does not imply that there is no DNA or RNA from other sources. The other source DNA may, for example, comprise DNA from a yeast or bacterial genome, or a cloning vector such as pUC19. This term distinguishes from naturally occurring events, such as viral infection, or tumor type growths, in which the level of one mRNA may be naturally increased relative to other species of mRNA. That is, the term is meant to cover only those situations in which a person has intervened to elevate the proportion of the desired nucleic acid.

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It is also advantageous for some purposes that a nucleotide sequence be in purified form. The term "purified" in reference to nucleic acid does not require absolute purity (such as a homogeneous preparation). Instead, it represents an indication that the sequence is relatively more pure than in the natural environment (compared to the natural level, this level should be at least 2-5 fold greater, e.g., in terms of mg/mL). Individual clones isolated from a cDNA library may be purified to electrophoretic homogeneity. The claimed DNA molecules obtained from these clones could be obtained directly from total DNA or from total RNA. The cDNA clones are not naturally occurring, but rather are preferably obtained via manipulation of a partially purified naturally occurring substance (messenger RNA). The construction of a cDNA library from mRNA involves the creation of a synthetic substance (cDNA) and pure individual cDNA clones can be isolated from the synthetic library by clonal selection of the cells carrying the cDNA library. Thus, the process which includes the construction of a cDNA library from mRNA and isolation of distinct cDNA clones yields an approximately 106-fold purification of the native message. Thus, purification of at least one order of magnitude, preferably two or three orders, and more preferably four or five orders of magnitude is expressly contemplated.

The terms "isolated", "enriched", and "purified" as respect nucleic acids, above, may similarly be used to denote the relative purity and abundance of polypeptides (multimers of amino acids joined one to another by α-carboxyl:α-amino group (peptide) bonds). These, too, may be stored in, grown in, screened in, and selected from libraries using biochemical techniques familiar in the art. Such polypeptides may be natural, synthetic or chimeric and may be extracted using any of a variety of methods, such as antibody immunoprecipitation, other "tagging" techniques, conventional chromatography and/or electrophoretic methods. Some of the above utilize the corresponding nucleic acid sequence.

18

As indicated above, aspects and embodiments of the invention are not limited to entire genes and proteins. The invention also provides and utilizes fragments and portions thereof, preferably those which are "active" in the inhibitory sense described above. Such peptides or oligopeptides and oligo or polynucleotides have preferred lengths as specified above for nucleic acid and amino acid sequences from phage; corresponding recombinant constructs can be made to express the encoded same. Also included are homologous sequences and fragments thereof.

Nucleic acid sequences of the present invention can be isolated using a method similar to those described herein or other methods known to those skilled in the art. In addition, such nucleic acid sequences can be chemically synthesized by well-known methods. Also, by having particular phage ORFs, e.g., the phage ORFs identified herein (e.g., anti-bacterial ORFs of the present invention, portions thereof, or oligonucleotides derived therefrom as described), other antimicrobial sequences from other bacteriophage sources can be identified and isolated using methods described here or other methods, including methods utilizing nucleic acid hybridization and/or computer-based sequence alignment methods.

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The invention also provides bacteriophage antimicrobial DNA segments from other phages based on nucleic acids and sequences hybridizing to the presently identified inhibitory ORF under high stringency conditions or sequences that are highly homologous. The bacteriophage segment from a specific phage, e.g., an antimicrobial DNA segment, can be used to identify a related segment from another unrelated phage based on stringent conditions of hybridization or on being a homolog based on nucleic acid and/or amino acid sequence comparisons. As with identified inhibitory sequences, such homologous coding sequences and products can be used as antimicrobials, to construct active portions or derivatives, to construct peptidomimetics, and to identify bacterial targets.

The nucleotide and amino acid sequences identified herein are believed to be correct, however, certain sequences may contain a small percentage of errors, e.g., 1-5%. In the event that any of the sequences have errors, the corrected sequences can be readily provided by one skilled in the art using routine methods. For example, the nucleotide sequences can be confirmed or corrected by obtaining and culturing the relevant phage, and purifying phage genomic nucleic acids: A region or regions of interest can be amplified, e.g., by PCR from the appropriate genomic template, using primers based on the described sequence. The amplified regions can then be sequenced using any of the available methods (e.g., a dideoxy termination method).

19

This can be done redundantly to provide the corrected sequence or to confirm that the described sequence is correct. Alternatively, a particular sequence or sequences can be identified and isolated as an insert or inserts in a phage genomic library and isolated, amplified, and sequenced by standard methods. Confirmation or correction of a nucleotide sequence for a phage gene provides an amino acid sequence of the encoded product by merely reading off the amino acid sequence according to the normal codon relationships and/or expressed in a standard expression system and the polypeptide product sequenced by standard techniques. The sequences described herein thus provide unique identification of the corresponding genes, coding sequences, and other sequences, allowing those sequences to be used in the various aspects of the present invention.

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In other aspects, the invention provides recombinant vectors and cells harboring at least one of the phage ORFs or portion thereof, or bacterial target sequences described herein. As understood by those skilled in the art, vectors may be provided in different forms, including, for example, plasmids, cosmids, and virusbased vectors. See, e.g., Maniatis, T. et al. (1989) Molecular Cloning: A Laboratory Manual, Cold Spring Harbor University Press, Cold Spring, N.Y.; See also, Ausubel, F.M. et al. (eds.) (1994) Current Protocols in Molecular Biology. John Wiley & Sons, Secaucus, N.J.

In preferred embodiments, the vectors will be expression vectors, preferably shuttle vectors that permit cloning, replication, and expression within bacteria. An "expression vector" is one having regulatory nucleotide sequences containing transcriptional and translational regulatory information that controls expression of the nucleotide sequence in a host cell. Preferably the vector is constructed to allow amplification from vector sequences flanking an insert locus. In certain embodiments, the expression vectors may additionally or alternativley support expression, and/or replication in animal, plant and/or yeast cells due to the presence of suitable regulatory sequences, e.g., promoters, enhancers, 3' stabilizing sequences, primer sequences, etc. In preferred embodiments, the promoters are inducible and specific for the system in which expression is desired, e.g., bacteria, animal, plant, or yeast. The vectors may optionally encode a "tag" sequence or sequences to facilitate protein purification. Convenient restriction enzyme cloning sites and suitable selective marker(s) are also optionally included. Such selective markers can be, for example, antibiotic resistance markers or markers which supply an essential nutritive growth factor to an otherwise deficient mutant host, e.g., tryptophan, histidine, or leucine in the Yeast Two-Hybrid systems described below.

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The term "recombinant vector" relates to a single- or double-stranded circular nucleic acid molecule that can be transfected into cells and replicated within or independently of a cell genome. A circular double-stranded nucleic acid molecule can be cut and thereby linearized upon treatment with appropriate restriction enzymes. An assortment of nucleic acid vectors, restriction enzymes, and the knowledge of the nucleotide sequences cut by restriction enzymes are readily available to those skilled in the art. A nucleic acid molecule encoding a desired product can be inserted into a vector by cutting the vector with restriction enzymes and ligating the two pieces together. Preferably the vector is an expression vector, *e.g.*, a shuttle expression vector as described above.

By "recombinant cell" is meant a cell possessing introduced or engineered nucleic acid sequences, e.g., as described above. The sequence may be in the form of or part of a vector or may be integrated into the host cell genome. Preferably the cell is a bacterial cell.

In another aspect, the invention also provides methods for identifying and/or screening compounds "active on" at least one bacterial target of a bacteriophage inhibitor protein or RNA. Preferred embodiments involve contacting such a bacterial target or targets (e.g., bacterial target proteins) with a test compound, and determining whether the compound binds to or reduces the level of activity of the bacterial target (e.g., a bacterial target protein). Preferably this is done either in vivo (i.e., in a cell-based assay) or in vitro, e.g., in a cell-free system under approximately physiological conditions.

The compounds that can be used may be large or small, synthetic or natural, organic or inorganic, proteinaceous or non-proteinaceous. In preferred embodiments, the compound is a peptidomimetic, as described herein, a bacteriophage inhibitor protein or fragment or derivative thereof, preferably an "active portion", or a small molecule.

In preferred embodiments, the bacterial target is a target of a phage ORF identified herein, e.g., S. aureus phage 44AHJD ORF 1, 9, or 12, Streptococcus pneumoniae phage Dp-1 ORF 001, 002, 004, 008, 010, 013, 016, 021, 029, 030, 038, or 041, or Enterococcus sp. phage 182 ORF 002, 008, or 014.

In particular embodiments, the methods include the identification of bacterial targets or the site of action of an inhibitor on a bacterial target as described above or otherwise described herein.

In embodiments involving binding assays, preferably binding is to a fragment or portion of a bacterial target protein, where the fragment includes less than 90%, 80%, 70%, 60%, 50%, 40%, or 30% of an intact bacterial target protein. Preferably,

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the at least one bacterial target includes a plurality of different targets of bacteriophage inhibitor proteins, preferably a plurality of different targets. The plurality of targets can be in or from a plurality of different bacteria, but preferably is from a single bacterial species.

A "method of screening" refers to a method for evaluating a relevant activity or property of a large plurality of compounds (e.g., a bacteria-inhibiting activity), rather than just one or a few compounds. For example, a method of screening can be used to conveniently test at least 100, more preferably at least 1000, still more preferably at least 10,000, and most preferably at least 100,000 different compounds, or even more.

In the context of this invention, the term "small molecule" refers to compounds having molecular mass of less than 2000 Daltons, preferably less than 1500, still more preferably less than 1000, and most preferably less than 600 Daltons. Preferably but not necessarily, a small molecule is not an oligopeptide.

In a related aspect or in preferred embodiments, the invention provides a method of screening for potential antibacterial agents by determining whether any of a plurality of compounds, preferably a plurality of small molecules, is active on at least one target of a bacteriophage inhibitor protein or RNA. Preferred embodiments include those described for the above aspect, including embodiments which involve determining whether one or more test compounds bind to or reduce the level of activity of a bacterial target, and embodiments which utilize a plurality of different targets as described above.

The identification of bacteria-inhibiting phage ORFs and their encoded products also provides a method for identifying an active portion of such an encoded product. This also provides a method for identifying a potential antibacterial agent by identifying such an active portion of a phage ORF or ORF product. In preferred embodiments, the identification of an active portion involves one or more of mutational analysis, deletion analysis, or analysis of fragments of such products. The method can also include determination of a 3-dimensional structure of an active portion, such as by analysis of crystal diffraction patterns. In further embodiments, the method involves constructing or synthesizing a peptidomimetic compound, where the structure of the peptidomimetic compound corresponds to the structure of the active portion. In this context, "corresponds" means that the peptidomimetic compound structure has sufficient similarities to the structure of the active portion that the peptidomimetic will interact with the same molecule as the phage protein and preferably will elicit at least one cellular response in common which relates to the inhibition of the cell by the phage protein.

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In preferred embodiments, the ORF or ORF product is or is derived or obtained from *S. aureus* phage 44AHJD ORF 1, 9, or 12, *Streptococcus pneumoniae* phage Dp-1 ORF 001, 002, 004, 008, 010, 013, 016, 021, 029, 030, 038, or 041, or *Enterococcus* sp. phage 182 ORF 002, 008, or 014 or product thereof.

The methods for identifying or screening for compounds or agents active on a bacterial target of a phage-encoded inhibitor can also involve identification of a phage-specific site of action on the target.

Preferably in the methods for identifying or screening for compounds active on such a bacterial target, the target is uncharacterized; the target is from an uncharacterized bacterium from Table 1; the site of action is a phage-specific site of action.

Further embodiments include the identification of inhibitor phage ORFs and bacterial targets as in aspects above.

An "active portion" as used herein denotes an epitope, a catalytic or regulatory domain, or a fragment of a bacteriophage inhibitor protein that is responsible for, or a significant factor in, bacterial target inhibition. The active portion preferably may be removed from its contiguous sequences and, in isolation, still effect inhibition.

By "mimetic" is meant a compound structurally and functionally related to a reference compound that can be natural, synthetic, or chimeric. In terms of the present invention, a "peptidomimetic," for example, is a compound that mimics the activity-related aspects of the 3-dimensional structure of a peptide or polyeptide in a non-peptide compound, for example mimics the structure of a peptide or active portion of a phage- or bacterial ORF-encoded polypeptide.

A related aspect provides a method for inhibiting a bacterial cell by contacting the bacterial cell with a compound active on a bacterial target of a bacteriophage inhibitor protein or RNA, where the target was uncharacterized. In preferred embodiments, the compound is such a protein, or a fragment or derivative thereof; a structural mimetic, e.g., a peptidomimetic, of such a protein or fragment; a small molecule; the contacting is performed in vitro, the contacting is performed in vivo in an infected or at risk organism, e.g., an animal such as a mammal or bird, for example, a human, or other mammal described herein; the bacterium is selected from a genus and/or species listed in Table 1; the bacteriophage inhibitor protein is uncharacterized; the bacteriophage inhibitor protein is from an uncharacterized phage listed in Table 1; the phage inhibitor protein is from one of S. aureus phage 44AHJD ORF 1, 9, or 12, Streptococcus pneumoniae phage Dp-1 ORF 001, 002, 004, 008, 010, 013, 016, 021, 029, 030, 038, or 041, or Enterococcus sp. phage 182 ORF 002, 008, or 014.

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In the context of targets in this invention, the term "uncharacterized" means that the target was not recognized as an appropriate target for an antibacterial agent prior to the filing of the present application or alternatively prior to the present invention. Such lack of recognition can include, for example, situations where the target and/or a nucleotide sequence encoding the target were unknown, situations where the target was known, but where it had not been identified as an appropriate target or as an essential cellular component, and situations where the target was known as essential but had not been recognized as an appropriate target due to a belief that the target would be inaccessible or otherwise that contacting the cell with a compound active on the target in vitro would be ineffective in cellular inhibition, or ineffective in treatment of an infection. Methods described herein utilizing bacterial targets, e.g., for inhibiting bacteria or treating bacterial infections, can also utilize "uncharacterized target sites", meaning that the target has been previously recognized as an appropriate target for an antibacterial agent, but where an agent or inhibitor of the invention is used which acts at a different site than that at which the previously utilized antibacterial agent, i.e., a phage-specific site. Preferably the phage-specific site has different functional characteristics from the previously utilized site. In the context of targets or target sites, the term "phage-specific" indicates that the target or site is utilized by at least one bacteriophage as an inhibitory target and is different from previously identified targets or target sites.

In the context of this invention, the term "bacteriophage inhibitor protein" refers to a protein encoded by a bacteriophage nucleic acid sequence which inhibits bacterial function in a host bacterium. Thus, it is a bacteria-inhibiting phage product.

In the context of this invention, the phrase "contacting the bacterial cell with a compound active on a bacterial target of a bacteriophage inhibitor protein" or equivalent phrases refer to contacting with an isolated, purified, or enriched compound or a composition including such a compound, but specifically does not rely on contacting the bacterial cell with an intact phage which encodes the compound. Preferably no intact phage are involved in the contacting.

Related aspects provide methods for prophylactic or therapeutic treatment of a bacterial infection by administering to an infected, challenged or at risk organism a therapeutically or prophylactically effective amount of a compound active on a target of a bacteriophage inhibitor protein or RNA, or as described for the previous aspect. Preferably the bacterium involved in the infection or risk of infection produces the identified target of the bacteriophage inhibitor protein or alternatively produces a homologous target compound. In preferred embodiments, the host organism is a plant or animal, preferably a mammal or bird, and more preferably, a human or other

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mammal described herein. Preferred embodiments include, without limitation, those as described for the preceding aspect.

Compounds useful for the methods of inhibiting, methods of treating, and pharmaceutical compositions can include novel compounds, but can also include compounds which had previously been identified for a purpose other than inhibition of bacteria. Such compounds can be utilized as described and can be included in pharmaceutical compositions.

In preferred embodiments of this and other aspects of the invention utilizing bacterial target sequences of a bacteriiophage inhibitory ORF product, the target sequence is encoded by a *Staphylococcus* nucleic acid coding sequence, preferably *S. aureus*, a *Streptococcus* nucleic acid coding sequence, preferably *Streptococcus* pneumoniae, or *Enterococcus* nucleic acid coding sequence. Possible target sequences are described herein by reference to sequence source sites.

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The amino acid sequence of a polypeptide target is readily provided by translating the corresponding coding region. For the sake of brevity, the sequences are not reproduced herein. For the sake of brevity, the sequences are described by reference to the GenBank entries instead of being written out in full herein. In cases where the TIGR or GenBank entry for a coding region is not complete, the complete sequence can be readily obtained by routine methods, e.g., by isolating a clone in a phage host genomic library, and sequencing the clone insert to provide the relevant coding region. The boundaries of the coding region can be identified by conventional sequence analysis and/or by expression in a bacterium in which the endogenous copy of the coding region has been inactivated and using subcloning to identify the functional start and stop codons for the coding region.

In the context of nucleic acid or amino acid sequences of this invention, the term "corresponding" indicates that the sequence is at least 95% identical, preferably at least 97% identical, and more preferably at least 99% identical to a sequence from the specified phage genome, a ribonucleotide equivalent, a degenerate equivalent (utilizing one or more degenerate codons), or a homologous sequence, where the homolog provides functionally equivalent biological function.

By "treatment" or "treating" is meant administering a compound or pharmaceutical composition for prophylactic and/or therapeutic purposes. The term "prophylactic treatment" refers to treating a patient or animal that is not yet infected but is susceptible to or otherwise at risk of a bacterial infection. The term "therapeutic treatment" refers to administering treatment to a patient already suffering from, infection.

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The term "bacterial infection" refers to the invasion of the host organism, animal or plant, by pathogenic bacteria. This includes the excessive growth of bacteria which are normally present in or on the body of the organism, but more generally, a bacterial infection can be any situation in which the presence of a bacterial population(s) is damaging to a host organism. Thus, for example, an organism suffers from a bacterial population when excessive numbers of a bacterial population are present in or on the organism's body, or when the effects of the presence of a bacterial population(s) is damaging to the cells, tissue, or organs of the organism.

The terms "administer", "administering", and "administration" refer to a method of giving a dosage of a compound or composition, e.g., an antibacterial pharmaceutical composition, to an organism. Where the organism is a mammal, the method is, e.g., topical, oral, intravenous, transdermal, intraperitoneal, intramuscular, or intrathecal. The preferred method of administration can vary depending on various factors, e.g., the components of the pharmaceutical composition, the site of the potential or actual bacterial infection, the bacterium involved, and the infection severity.

The term "mammal" has its usual biological meaning referring to any organism of the Class Mammalia of higher vertebrates that nourish their young with milk secreted by mammary glands, e.g., mouse, rat, and, in particular, human, bovine, sheep, swine, dog, and cat.

In the context of treating a bacterial infection a "therapeutically effective amount" or "pharmaceutically effective amount" indicates an amount of an antibacterial agent, e.g., as disclosed for this invention, which has a therapeutic effect. This generally refers to the inhibition, to some extent, of the normal cellular functioning of bacterial cells that renders or contributes to bacterial infection.

The dose of antibacterial agent that is useful as a treatment is a "therapeutically effective amount." Thus, as used herein, a therapeutically effective amount means an amount of an antibacterial agent that produces the desired therapeutic effect as judged by clinical trial results and/or animal models. This amount can be routinely determined by one skilled in the art and will vary depending on several factors, such as the particular bacterial strain involved and the particular antibacterial agent used.

In connection with claims to methods of inhibiting bacteria and therapeutic or prophylactic treatments, "a compound active on a target of a bacteriophage inhibitor protein" or terms of equivalent meaning differ from administration of or contact with an intact phage naturally encoding the full-length inhibitor compound. While an intact phage may conceivably be incorporated in the present methods, the method at

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least includes the use of an active compound as specified different from a full length inhibitor protein naturally encoded by a bacteriophage and/or a delivery or contacting method different from administration of or contact with an intact phage encoding the full-length protein. Similarly, pharmaceutical compositions described herein at least include an active compound different from a full-length inhibitor protein naturally encoded by a bacteriophage or such a full-length protein is provided in the composition in a form different from being encoded by an intact phage. Preferably the methods and compositions do not include an intact phage.

In accord with the above aspects, the invention also provides antibacterial agents and compounds active on bacterial targets of bacteriophage inhibitor proteins or RNAs, where the target was uncharacterized as indicated above. As previously indicated, such active compounds include both novel compounds and compounds which had previously been identified for a purpose other than inhibition of bacteria. Such previously identified biologically active compounds can be used in embodiments of the above methods of inhibiting and treating. In preferred embodiments, the targets, bacteriophage, and active compound are as described herein for methods of inhibiting and methods of treating. Preferably the agent or compound is formulated in a pharmaceutical composition which includes a pharmaceutically acceptable carrier, excipient, or diluent. In addition, the invention provides agents, compounds, and pharmaceutical compositions where an active compound is active on an uncharacterized phage-specific site.

In preferred embodiments, the target is as described for embodiments of aspects above.

Likewise, the invention provides a method of making an antibacterial agent. The method involves identifying a target of a bacteriophage inhibitor polypeptide or protein or RNA, screening a plurality of compounds to identify a compound active on the target, and synthesizing the compound in an amount sufficient to provide a therapeutic effect when administered to an organism infected by a bacterium naturally producing the target. In preferred embodiments, the identification of the target and identification of active compounds include steps or methods and/or components as described above (or otherwise herein) for such identification. Likewise, the active compound can be as described above, including fragments and derivatives of phage inhibitor proteins, peptidomimetics, and small molecules. As recognized by those skilled in the art, peptides can be synthesized by expression systems and purified, or can be synthesized artificially. In preferred embodiments the inhibitory phage ORF products is from S. aureus phage 44AHJD ORF 1, 9, or 12, Streptococcus

pneumoniae phage Dp-1 ORF 001, 002, 004, 008, 010, 013, 016, 021, 029, 030, 038, or 041, or *Enterococcus* sp. phage 182 ORF 002, 008, or 014.

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As indicated above, sequence analysis of nucleotide and/or amino acid sequences can beneficially utilize computer analysis. Thus, in additional aspects the invention provides computer-related hardware and media and methods utilizing and incorporating sequence data from uncharacterized phage, e.g., uncharacterized phage listed in Table 1, preferably at least one of Staphylococcus aureus phage S. aureus phage 44AHJD ORF 1, 9, or 12, Streptococcus pneumoniae phage Dp-1 ORF 001, 002, 004, 008, 010, 013, 016, 021, 029, 030, 038, or 041, or Enterococcus sp. phage 182 ORF 002, 008, or 014, or 44 AHJD, Enterococcus sp. phage 182, or Streptococcus pneumoniae phage Dp-1. In general, such aspects can facilitate the above-described aspects. Various embodiments involve the analysis of genetic sequence and encoded products, as applied to the evaluating bacteriophage inhibitor ORFs and compounds and fragments related thereto. The various sequence analyses, as well as function analyses, can be used separately or in combination, as well as in preceding aspects and embodiments. Use in combination is often advantageous as the additional information allows more efficient prioritizing of phage ORFs for identification of those ORFs that provide bacteria-inhibiting function.

In one aspect, the invention provides a computer-readable device which includes at least one recorded amino acid or nucleotide sequence corresponding to one of the specified phage and a sequence analysis program for analyzing a nucleotide and/or amino acid sequence. The device is arranged such that the sequence information can be retrieved and analyzed using the analysis program. The analysis can identify, for example, homologous sequences or the indicated %s of the phage genome and structural motifs. Preferably the sequence includes at least 1 phage ORF or encoded product, more preferably at least 10%, 20%, 30%, 40%, 50%, 70%, 90%, or 100% of the genomic phage ORFs and/or equivalent cDNA, RNA, or amino acid sequences. Preferably the sequence or sequences in the device are recorded in a medium such as a floppy disk, a computer hard drive, an optical disk, computer random access memory (RAM), or magnetic tape. The program may also be recorded in such medium. The sequences can also include sequences from a plurality of different phage.

In this context, the term "corresponding" indicates that the sequence is at least 95% identical, preferably at least 97% identical, and more preferably at least 99% identical to a sequence from the specified phage genome, a ribonucleotide equivalent, a degenerate equivalent (utilizing one or more degenerate codons), or a homologous sequence, where the homolog provides functionally equivalent biological function.

Similarly, the invention provides a computer analysis system for identifying biologically important portions of a bacteriophage genome. The system includes a data storage medium, e.g., as identified above, which has recorded thereon a nucleotide sequence corresponding to at least a portion of at least one uncharacterized bacteriophage genome, a set of program instructions to allow searching of the sequence or sequences to analyze the sequence, and an output device where the portion includes at least the sequence length as specified in the preceding aspect. The output device is preferably a printer, a video display, or a recording medium. More one than one output device may be included. For each of the present computer-related asepcts, the bacteriophage are preferably selected from the uncharacterized phage listed in Table 1, more preferably from bacteriophage 77, 3A, 96, 44 AHJD (S. aureus), Dp-1 (Streptococcus pneumoniae), or 182 (Enterococcus).

In keeping with the computer device aspects, the invention also provides a method for identifying or characterizing a bacteriophage ORF by providing a computer-based system for analyzing nucleotide or amino acid sequences, e.g., as describe above. The system includes a data storage medium which has recorded a sequences or sequences as described for the above devices, a set of instructions as in the preceding aspect, and an output device as in the preceding aspect. The method further involves analyzing at least one sequence, and outputting the analysis results to at least one output device.

In preferred embodiments, the analysis identifies a sequence similarity or homology with a sequence or sequences selected from bacterial ORFs encoding products with related biological function; ORFs encoding known inhibitors; and essential bacterial ORFs. Preferably the analysis identifies a probable biological function based on identification of structural elements or characteristic or signature motifs of an encoded product or on sequence similarity or homology. Preferably the uncharacterized bacteriophage is from Table 1, more preferably at least one of bacteriophage 77, 3A, 96, 44 AHJD (S. aureus), Dp-1 (Streptococcus pneumoniae), or 182 (Enterococcus). In preferred embodiments, the method also involves determining at least a portion of the nucleotide sequence of at least one uncharacterized bacteriophage as indicated, and recording that sequence on data storage medium of the computer-based system. In preferred embodiments, the analysis identifies a sequence similarity of homology with a S. aureus phage 44AHJD ORF 1, 9, or 12, Streptococcus pneumoniae phage Dp-1 ORF 001, 002, 004, 008, 010, 013, 016, 021, 029, 030, 038, or 041, or Enterococcus sp. phage 182 ORF 002, 008, or 014.

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As used in the claims to describe the various inventive aspects and embodiments, "comprising" means including, but not limited to, whatever follows the word "comprising". Thus, use of the term "comprising" indicates that the listed elements are required or mandatory, but that other elements are optional and may or may not be present. By "consisting of" is meant including, and limited to, whatever follows the phrase "consisting of". Thus, the phrase "consisting of" indicates that the listed elements are required or mandatory, and that no other elements may be present. By "consisting essentially of" is meant including any elements listed after the phrase, and limited to other elements that do not interfere with or contribute to the activity or action specified in the disclosure for the listed elements. Thus, the phrase "consisting essentially of" indicates that the listed elements are required or mandatory, but that other elements are optional and may or may not be present depending upon whether or not they affect the activity or action of the listed elements.

Further embodiments will be apparent from the following Detailed Description and from the claims.

# **BRIEF DESCRIPTION OF THE DRAWINGS**

FIGURE 1A and 1B are flow schematics showing the manipulations used to convert pT0021, an arsenite inducible vector containing the luciferase gene, into pTHA or pTM, two *ars* inducible vectors. Vector pTHA contains BamH I, Sal I, and Hind III cloning sites and a downstream HA epitope tag. Vector pTM contains Bam HI and Hind III cloning sites and no HA epitope tag.

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FIGURE 2 is a schematic representation of the cloning steps involved to place the DNA segments of any of ORFs 17/ 19/ 43/ 102/104/182 or other sequences into pTHA to assess inhibitory potential. For subcloning into pTM or pT0021, Individual ORFs were amplified by the PCR using oligonucleotides targeting the ATG and stop codons of the ORFs. Using this strategy, Bam HI and Hind III sites were positioned immediately upstream or downstream, respectively of the start and stop codons of each ORF. Following digestion with Bam HI and Hind III, the PCR fragments were subcloned into the same sites of pT0021 or pTM. Clones were verified by PCR and direct sequencing.

FIGURE 3 shows a schematic representation of the functional assays used to characterize the bactericidal and bacteriostatic potential of all predicted ORFs (>33 amino acids) encoded by bacteriophage 77. Fig. 3A) Functional assay on semi-solid support media. Fig. 3B) Functional assay in liquid culture.

FIGURE 4A, B, and C is a bar graph showing the results of a screen in liquid media to assess bacteriostatic or bactericidal activity of 93 predicted ORFs (>33 amino acids) encoded by bacteriophage 77. Growth inhibition assays were performed as detailed in the Detailed Description. The relative growth of Staphylococcus aureus transformants harboring a given bacteriophage 77 ORF (identified on the bottom of the graph), in the absence or presence of arsenite, is plotted relative to growth of a Staphylococcus aureus transformant containing ORF 5, a non-toxic bacteriophage 77 ORF (which is set at 100%). Each bar represents the average obtained from three Staph A transformants grown in duplicate. Bacteriophage 77 ORFs showing significant growth inhibition consist of ORFs 17, 19, 102, 104, and 182.

FIGURE 5 shows a block diagram of major components of a general purpose computer.

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FIGURE 6 shows an ORF map for *Streptococcus pneumoniae* bacteriophage Dp-1 showing the ORF identifiers, genomic locations, and orientations of the 85 identified ORFs that were found to have ribosomal binding sites and thus are expected to be expressed.

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FIGURE 7 shows a schematic representation of the arsenite-inducible expression system present in a shuttle vector designed to express individual *Streptococcus* bacteriophage Dp-1 ORFs in *Streptococcus*. Various modifications can be readily made to such a vector, or other vectors can be readily constructed to provide inducible expression of ORFs in a particular host bacterium using well-known techniques.

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# **DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS**

The invention may be more clearly understood from the following description.

The tables will first be briefly described.

Table 1 is a listing of a large number of available bacteriophage that can be readily obtained and used in the present invention.

Table 2 shows the complete nucleotide sequence of the genome of Staphylococcus aureus bacteriophage 77.

Table 3 shows a list of all the ORFs from Bacteriophage 77 that were screened in the functional assay to identify those with anti-microbial activity.

Table 4 shows the predicted nucleotide sequence, predicted amino acid sequence, and physiochemical parameters of ORF 17/ 19/ 43/ 102/ 104/ 182]. These include the primary amino acid sequence of the predicted protein, the average molecular weight, amino acid composition, theoretical pI, hydrophobicity map, and predicted secondary structure map.

Table 5 shows homology search results. BLAST analysis was performed with ORFs 17/ 19/ 43/ 102/ 104/ 182 against NCBI non-redundant nucleotide and Swissprot databases. The results of this search indicate that: I) ORF 17 has no significant homology to any gene in the NCBI non-redundant nucleotide database, II) ORF 19 has significant homology to one gene in the NCBI non-redundant nucleotide database - the gene encoding ORF 59 of bacteriophage phi PVL, III) ORF 43 has significant homology to one gene in the NCBI non-redundant nucleotide database - the gene encoding ORF 39 of phi PVL, IV) ORF 102 has significant homology to one gene in the NCBI non-redundant nucleotide database - the gene encoding ORF 38 of phi PVL, V) ORF 104 has no significant homology to any gene in the NCBI non-redundant nucleotide database, VI) ORF 182 has significant homology to one gene in the NCBI non-redundant nucleotide database - the gene encoding ORF 39 of phi PVL.

Table 6 is a table from Alberts et al., MOLECULAR BIOLOGY OF THE CELL 3<sup>rd</sup> ed., showing the redundancy of the "universal" genetic code.

Table 7 shows the complete nucleotide sequence of *Staphylococcus aureus* bacteriophage 3A.

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Table 8 is a listing of the ORFs identified in *Staphylococcus aureus* bacteriophage 3A.

Table 9 shows the complete nucleotide sequence of *Staphylococcus aureus* bacteriophage 96.

Table 10 is a listing of the ORFs identified in *Staphylococcus aureus* bacteriophage 96.

Table 11 is a listing of sequences deposited in the NCBI public database (GeneBank) for bacteriophage listed in Table 1.

Table 12 is a listing of phage which encode a known lysis function, including the identified lysis gene.

Table 13 is a listing of bacteriophage which encode holin genes, where holin genes encode proteins which form pores and eventually enable other enzymes to kill the host bacterium.

Table 14 is a listing of bacteriophage which encode kil genes.

Table 15 is a list of *Staphylococcus aureus* sequences identified by accession number which may include sequences from genes coding for target sequences for the phage 77-encoded antimicrobial proteins or peptides. The sequences were obtained by searching GenBank for listings.

Table 16 shows the nucleotide sequence of the genome of *Staphylococcus* aureus phage 44 AHJD.

Table 17 lists and shows the sequence position of the 73 ORFs predicted to be encoded by *Staphylococcus aureus* bacteriophage 44 AHJD that are greater than 33 amino acids.

Table 18 shows the ORF sequences and putative amino acid sequences for the Staphylococcus aureus bacteriophage 44AHJD ORFs greater than 33 amino acids.

Table 19 shows the similarities in sequence identified between predicted Staphylococcus aureus bacteriophage 44 AHJD ORFs and sequences present in public databases.

Table 20 shows the homology alignments between predicted *Staphylococcus* aureus bacteriophage 44AHJD ORFs and the corresponding protein sequences present in public sequence databases.

Table 21 shows the complete nucleotide sequence of the genome of *Enterococcus* bacteriophage 182.

Table 22 lists and shows the sequence position of the 80 ORFs identified in bacteriophage 182 and that are greater than 33 amino acids.

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Table 23 shows the nucleotide and predicted amino acid sequence of all 80 ORFs identified in bacteriophage 182.

Table 24 shows the similarities identified to date in sequence between Enterococcus phage 182 ORFs greater than 33 amino acids and sequences present in public sequence databases.

Table 25 shows the predicted amino acid sequence as well as the predicted secondary structures map for two *Enterococcus* bacteriophage 182 ORFs.

Table 26 shows the homology alignments between predicted *Enterococcus* bacteriophage 182 ORFs and the corresponding protein sequences present in public sequence databases.

Table 27 list *Enterococcus* sequences listed in GenBank providing possible Enterococcal target sequences for inhibitory *Enterococcus* bacteriophage 182 ORFs and other compounds with antibacterial activity.

Table 28 shows the complete nucleotide sequence of the genome of *Streptococcus* bacteriophage Dp-1.

Table 29 lists and shows sequence position of the 273 ORFs identified in Pneumococcal bacteriophage Dp-1 that are greater than 33 amino acids, 85 of which are predicted to be expressed in Dp-1 as having a ribosomal binding site. That set of 85 ORFs is shown in the attached drawings.

Table 30 shows the nucleotide and predicted amino acid sequence of all 273 ORFs identified in bacteriophage Dp-1 that are identified as being expressed.

Table 31 shows the similarities identified in sequence between *Streptococcus* phage Dp-1 ORFs greater than 33 amino acids and sequences present in public sequence databases.

Table 32 shows the 4731 bp sequence of Dp-1 published by Sheehan et al., 1997).

Table 33 lists *Streptococcus pneumoniae* sequences listed in GenBank providing possible target sequences for inhibitory *Streptococcus pneumoniae* bacteriophage Dp-1 ORFs and other compounds with antibacterial activity

# Background:

As indicated above, the present invention is concerned, in part, with the use of bacteriophage coding sequences and the encoded polypeptides or RNA transcripts to identify bacterial targets for potential new antibacterial agents. Thus, the invention concerns the selection of relevant bacteria. Particularly relevant bacteria are those which are pathogens of a complex organism such as an animal, e.g., mammals,

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reptiles, and birds, and plants. Examples include *Stapylococcus aureus*, *Enterococcus* species, and *Streptococcus pneumoniae*. However, the invention can be applied to any bacterium (whether pathogenic or not) for which bacteriophage are available or which are found to have cellular components closely homologous to components targeted by phage of another bacterium.

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Thus, the invention also concerns the bacteriophage which can infect a selected bacterium. Identification of ORFs or products from the phage which inhibit the host bacterium both provides an inhibitor compound and allows identification of the bacterial target affected by the phage-encoded inhibitor. Such targets are thus identified as potential targets for development of other antibacterial agents or inhibitors and the use of those targets to inhibit those bacteria. As indicated above, even if such a target is not initially identified in a particular bacterium, such a target can still be identified if a homologous target is identified in another bacterium. Usually, but not necessarily, such another bacterium would be a genetically closely related bacterium. Indeed, in some cases, a phage-encoded inhibitor can also inhibit such a homologous bacterial cellular component.

The demonstration that bacteriophage have adapted to inhibiting a host bacterium by acting on a particular cellular component or target provides a strong indication that that component is an appropriate target for developing and using antibacterial agents, e.g., in therapeutic treatments. Thus, the present invention provides additional guidance over mere identification of bacterial essential genes, as the present invention also provides an indication of accessability of the target to an inhibitor, and an indication that the target is sufficiently stable over time (e.g., not subject to high rates of mutation) as phage acting on that target were able to develop and persist. Thus, the present invention identifies a subset of essential cellular components which are particularly likely to be appropriate targets for development of antibacterial agents.

The invention also, therefore, concerns the development or identification of inhibitors of bacteria, in addition to the phage-encoded inhibitory proteins (or RNA transcripts), which are active on the targets of bacteriophage-encoded inhibitors. As described herein, such inhibitors can be of a variety of different types, but are preferably small molecules.

The following description provides preferred methods for use in the various aspects of the invention. However, as those skilled in the art will readily recognize, other approaches can be used to obtain and process relevant information. Thus the invention is not limited to the specifically described methods. In addition, the following description provides a set of steps in a particular order. That series of steps

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describes the overall development involved in the present invention. However, it is clear that individual steps or portions of steps may be usefully practiced separately, and, further, that certain steps may be performed in a different order or even bypassed if appropriate information is already available or is provided by other sources or methods.

# Selecting and Growing Phage, and Isolating DNA

Conceptually, the first step involves selecting bacterial hosts of interest. Preferably, but not necessarily, such hosts will be pathogens of clinical importance. Alternatively, because bacteria all share certain fundamental metabolic and structural features, these features can be targeted for study in one strain, for example a nonpathogenic one, and extrapolated to similarly succeed in pathogenic ones. Nonpathogenic strains may also exhibit initial advantages in being not only less dangerous, but also, for example, in having better growth and culturing characteristics and/or better developed molecular biology techniques and reagents. Consequently, advantageously the invention provides the ability target virtually any bacteria, but preferably pathogenic bacteria, with antimicrobial compounds designed and/or developed using bacteriophage inhibitory proteins and peptides from phage with non-pathogenic and/or pathogenic hosts.

We have selected Staphylococcus aureus, Streptococcus pneumoniae, various Enterococci, and Pseudomonas aeruginosa as initial exemplary pathogens. These bacteria are a major cause of morbidity and mortality in hospital-based infections, and the appearance of antibiotics resistance in all three organisms makes it increasingly difficult to treat benign infections involving these organisms. Such infections can include, for example, otitis media, sinusitis, and skin, and airway infections (Neu, H.C. (1992). Science 257, 1064-1073). However, the approach described below is clearly applicable to any human bacterial pathogens including but not restricted to Mycobacterium tuberculosis, Nesseria gonorrhoeae, Haemophilus influenza, Acinobacter, Escherichia coli, Shigella dysenteria, Streptococcus pyogenes, Helicobacter pylori, and Mycoplasma species. This invention can also be applied to the discovery of anti-bacterial compounds directed against pathogens of animals other than humans, for example, sheep, cattle, swine, dogs, cats, birds, and reptiles. Similarly, the invention is not limited to animals, but also applies to plants and plant pathogens.

In general, the bacteria are grown according to standard methodologies - employed in the art, including solid, semi-solid or liquid culturing, which procedures can be found in or extrapolated from standard sources such as Maloy, S.R., Stewart,

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V.J., and Taylor, R.K. Genetic Analysis of Pathogenic Bacteria (1996) Cold Spring Harbor Laboratory Press, or Maniatis, T. et al. (1989) Molecular Cloning: A Laboratory Manual, Cold Spring Harbor University Press, Cold Spring, N.Y.; or Ausubel, F.M. et al. (1994) Current Protocols in Molecular Biology. John Wiley & Sons, Secaucus, N.J. Culture conditions are selected which are adapted to the particular bacterium generally using culture conditions known in the art as appropriate, or adaptations of those conditions.

Nucleic acids within these bacteria can be routinely extracted through common procedures such as described in the above-referenced manuals and as generally known to those skilled in the art. Those nucleic acid stocks can then be used to practice the other inventive aspects described below.

# Selection and Growth of Bacteriophage, and Isolation of DNA

The second step involves assembling a group of bacteriophages (phage collection) for one or more of the targeted bacterial hosts. While the invention can be utilized with a single bacteriophage for a pathogen or other bacterium, it is preferable to utilize a plurality of phage for each bacterium, as comparisons between a plurality of such phage provides useful additional information. Non-limiting examples of phage and sources for some of the above-mentioned pathogenic bacteria are found in Table 1. The criteria used to select such phages is that they are infectious for the microbe targeted, and replicate in, lyse, or otherwise inhibit growth of the bacterium in a measurable fashion. These phages can be very different from one another (representing different families), as judged by criteria such as morphology (head, tail, plate, etc.), and similarity of genome nucleotide sequence (cross-hybridization). Since such diverse bacteriophages are expected to block bacterial host metabolism and ultimately inhibit by a variety of mechanisms, their combined study will lead to the identification of different mechanisms by which the phages independently inhibit bacterial targets. Examples include degradation of host DNA (Parson K.A., and Snustad, D.P. (1975). J. Virol. 15, 221-444) and inhibition of host RNA transcription (Severinova, E., Severinov, K. and Darst, S.A. (1998). J.Mol. Biol. 279, 9-18). This, in turn, yields novel information on phage proteins that can inhibit the targeted microbe. As explained below, this 1) forms the basis of novel drug discovery efforts based on knowledge of the primary amino acid sequence of the phage inhibitor protein (e.g., peptide fragments or peptidomimetics) and/or 2) leads to the identification of bacterial biochemical pathways, the proteins of which are essential or significant for survival of the targeted microbe, and which enzymatic steps or

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chemical reactions can be targeted by classical drug discovery methods using molecular inhibitors, for example, small molecule inhibitors.

Bacteriophage are generally either of two types, lytic or filamentous, meaning they either outright destroy their host and seek out new hosts after replication, or else continuously propogate and extrude progeny phage from the same host without destroying it. Regardless of the phage life cycle and type, preferred embodiments incorporate phage which impede cell growth in measurable fashion and preferably stop cell growth. To this end, lytic phage are preferred, although certain nonlytic species may also suffice, e.g., if sufficiently bacteriostatic.

Various procedures that are commonly understood by those of skill in the art can be routinely employed to grow, isolate, and purify phage. Such procedures are exemplified by those found in such common laboratory aids such as Maloy, S.R., Stewart, V.J., and Taylor, R.K. Genetic Analysis of Pathogenic Bacteria (1996) Cold Spring Harbor Laboratory Press; Maniatis, T. et al. (1989) Molecular Cloning: A Laboratory Manual, Cold Spring Harbor University Press, Cold Spring, N.Y.; and Ausubel, F.M. et al. (eds.) (1994) Current Protocols in Molecular Biology. John Wiley & Sons, Secaucus, N.J. The techniques generally involve the culturing of infected bacterial cells that are lysed naturally and/or chemically assisted, for example, by the use of an organic solvent such as chloroform that destroys the host cells thereby liberating the phage within. Following this, the cellular debris is centrifuged away from the supernatant containing the phage particles, and the phage then subsequently and selectively precipitated out of the supernatant using various methods usually employing the use of alcohols and/or other chemical compounds such as polyethylene glycol (PEG). The resulting phage can be further purified using various density gradient/centrifugation methodologies. The resulting phage are then chemically lysed, thereby releasing their nucleic acids that can be conveniently precipitated out of the supernatant to yield a viral nucleic acid supply of the phage of interest.

Exemplary bacteriophage are indicated in Table 1, along with sources where those phage may be obtained.

Exemplary bacteria include the reference bacteria for the identified bacteriophage, available from the same sources.

## Characterizing Bacteriophage Genomes for ORFs

The third step involves systematically characterizing the genetic information contained in the phage genome. Within this genetic information is the sequence of all RNAs and proteins encoded by the phage, including those that are essential or

38

instrumental in inhibiting their host. This characterization is preferably done in a systematic fashion. For example, this can be done by first isolating high molecular weight genomic DNA from the phage using standard bacterial lysis methods, followed by phage purification using density gradient ultracentrifugation, and extraction of nucleic acid from the purified phage preparation. The high molecular weight DNA is then analyzed to determine its size and to evaluate a proper strategy for its sequencing. The DNA is broken down into smaller size fragments by sonication or partial digestion with frequently cutting restriction enzymes such as Sau3A to yield predominantly 1 to 2 kilobase length DNA, which DNA can then be resolved by gel electrophoresis followed by extraction from the gel.

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The ends of the fragments are enzymatically treated to render them suitable for cloning and the pools of fragments are cloned in a bacterial plasmid to generate a library of the phage genome. Several hundred of these random DNA fragments contained in the plasmid vector are isolated as clones after introduction into an appropriate bacterium, usually *Escherichia coli*. They are then individually expanded in culture and the DNA from each individual clone is purified. The nucleotide sequences of the inserts of these clones are determined by standard automated or manual methods, using oligonucleotide primers located on either side of the cloning site to direct polymerase mediated sequencing (e.g., the Sanger sequencing method or a modification of that method). Other sequencing methods can also be used.

The sequence of individual clones is then deposited in a computer, and specific software programs (for example, Sequencher<sup>TM</sup>, Gene Codes Corp.) are used to look for overlap between the various sequences, resulting in ordering of contig sequences and ultimately providing the complete sequence of the entire bacteriophage genome (one such example is given in Table 2 for *Staphylococcus aureus* bacteriophage 77; others are also provided herein). This complete nucleotide sequence is preferably determined with a redundancy of at least 3- to 5-fold (number of independent sequencing events covering the same region) in order to minimize sequencing errors.

Preferably, the bacterial strain used as a phage host should not possess any other innate plasmids, transposons, or other phage or incompatible sequences that would complicate or otherwise make the various manipulations and analyses more difficult.

Commercially available computer software programs are used to translate the nucleotide sequence of the phage to identify all protein sequences encoded by the phage (hereafter called open reading frames or ORFs). (Customized software can clearly also be used.) As phages are known to transcribe their genome into RNA from

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both strands, in both directions, and sometimes in more than one frame for the same sequence, this exercise is done for both strands and in all six possible reading frames. As evolutionary constraints have forced the phage to conserve all of its vital protein sequences in as small a genome as possible, it is straightforward to identify all the proteins encoded by the phage by simple examination of the 6 translation frames of the genome. Once these ORFs are identified, they are cataloged into a phage proteome database (Table 3 lists ORFs identified from phage 77; ORF lists are also provided for other exemplary phage). This analysis is preferably performed for each phage under study. The process of ORF identification can be varied depending on the desired results. For example, the minimum length for the putative encoded polypeptide can be varied, and/or putative coding regions that have an associated Shine-Dalgarno sequence can be selected. In the case of phage 77 ORFs, such parameter adjustment was performed and resulted in the identification of ORFs as listed herein. Different parameters had resulted in the identification of the ORFs listed in the preceding U.S. Provisional Application 60/110,992, filed December 3. 1998, which is hereby incorporated by reference in its entirety.

Exemplary phage 77 ORFs identified in that provisional application and as identified herein are shown in the following table:

ORF ID from 60/110,992	Genomic position	a.a. size	Start codon	ORF ID from 241/190	Genomic position	a.a. size	Start codon
77ORF016	2369-24024	251	TTG	77ORF017	23269-23982	237	ATG
77ORF019	39845-40501	218	ATA	77ORF019	39851-40501	216	ATG
77ORF050	29268-29564	98	ATG	77ORF182	29268-29564	98	ATG
77ORF050	29268-29564	98	ATG	77ORF043	29304-29564	86	ATG
77ORF067	34312-34551	79	CTG	77ORF104	34393-34551	52	ATG
770RF146	29051-29212	53	ATG	77ORF102	29051-29212	53	ATG

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## Identifying and Characterizing Inhibitory Phage ORFs

The fourth step entails identifying the phage protein or proteins or RNA transcripts that have the ability to inhibit their bacterial hosts. This can be accomplished, for example, by either or both of two non-mutually exclusive methods. The first method makes use of bioinformatics. Over the past few years, a large amount of nucleotide sequence information and corresponding translated products have become available through large genome sequencing projects for a variety of organisms including mammals, insects, plants, unicellular eukaryotes (yeast and fungi), as well as several bacterial genomes such as E. coli, Mycobacterium tuberculosis, Bacillus subtilis, Staphylococcus aureus and many others. Such sequences have been deposited in public databases (for example, non-redundant

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sequence database at GenBank and SwissProt protein sequence database) (http://www.ncbi.nlm.nih.gov)) and can be freely accessed to compare any specific query sequence to those present in such databases. For example, GenBank contains over 1.6 billion nucleotides corresponding to 2.3 million sequence records. Several computer programs and servers (e.g., TBLASTN) have been created to allow the rapid identification of homology between any given sequence from one organism to that of another present in such databases, and such programs are public and available free of charge.

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In addition, it has been well established that basic biochemical pathways can be conserved in very distant organisms (for example bacteria and man), and that the proteins performing the various enzymatic steps in these pathways are themselves conserved at the amino acid sequence level. Thus, proteins performing similar functions (e.g. DNA repair, RNA transcription, RNA translation) have frequently preserved key structural signatures, identifiable by similarities across regions of proteins (domains and motifs). The antimicrobials of the present invention will preferably target features and targets that are highly characteristic or conserved in microbes, and not higher organisms.

Most genomes encode individual proteins or groups of proteins that can be assembled into protein families that have been evolutionarily conserved. Therefore, similarity between a new query sequence and that of a member of a protein family (reference sequences from public databases) can immediately suggest a biochemical function for the novel query sequence, which in our case is a phage ORF.

The sequence homology between individual members of evolutionarily distant members of a protein family is usually not randomly distributed along the entire length of the sequence but is often clustered into "motifs" and "domains". These correspond to key three-dimensional folds that form key catalytic and/or regulatory structures that perform key biochemical function(s) for the group of proteins. Commercially available computer software programs can identify such motifs in a new query sequence, again providing functional information for the query sequence. Such structural and functional motifs have also been derived from the combined analysis of primary sequence databases (protein sequences) and protein structure databases (X-ray crystallography, nuclear magnetic resonance) using so-called "threading" methods (Rost B,l and Sander C. (1996). Ann. Rev. Biophy. Biomol. Struct. 25, 113-136).

Such motifs and folds are themselves deposited in public databases which can be directly accessed (for example, SwissProt database; 3D-ALI at EMBL, Heidelberg; PROSITE). This basic exercise leads to a structural homology map in which each of

41

the phage ORFs has been probed for such similarities, and where initial structural and functional hits are identified (selected examples of sequence homologies detected between individual ORFs from the genome of *Staphylococcus aureus* bacteriophage 77 and sequences deposited in public databases are shown in Table 5 for ORFs 17/19/43/102/104/182).

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This analysis can point out phage proteins with similarity to proteins from other phages (such as those for *E. coli*) playing an important role in the basic biochemical pathways of the phage (such as DNA replication, RNA transcription, tRNAs, coat protein and assembly). Selected examples of such proteins include integrase and capsid protein. Therefore, this analysis enables identification and elimination of non-essential ORFs as candidates for an inhibitor function, as well as the identification of (potentially) useful ones.

In addition, this analysis can point out specific ORFs as possible inhibitor ORFs. For example these ORFs may encode proteins or enzymes that alter bacterial cell structure, metabolism or physiology, and ultimately viability. Examples of such proteins present in the genome of *Staphylococcus aureus* bacteriophage 77 include orf14 (deoxyuridine triphosphatase from bacteriophage T5), and orf15 (sialidase). (These ORF identifications are as listed in provisional application 60/110,992.) Other examples include ORFs 9 and 12 of *S. aureus* phage 44 AHJD, which encode the putative lysis functions found in many bacteriophages – a "holin" and an "amidase".

In addition, it is well known that bacterial and eukaryotic viruses can usurp pathways from their host in order to use them to their advantage in blocking host cellular pathways upon infection. The phage can achieve this by 1) directly producing an inhibitor of a key host pathway (e.g. T7 gene 0.5 and 2), 2) directly producing a novel activity (e.g. T4 DNA polymerase), and 3) altering concentrations of cell components by producing similar functions (e.g. T4 transfer RNAs). The identification of sequence similarity between phage ORFs and bacterial host genome sequences will be highly indicative of such a mechanism. (Selected examples of such homologies are listed in Figure 4 of the provisional application 60/110,992 and include orf4 (homologous to autolysin), orf20 (hypothetical protein from Staphyloccus aureus.)) These ORFs can be analyzed by a standard biochemical approach to directly test their inhibitor functions (e.g., as described below).

Alternatively, a homology search may reveal that a given phage ORF is related to a protein present in the databases having an activity known to be inhibitory, ( $e.\overline{g}$ . inhibitor of host RNA polymerase by  $E.\ coli$  bacteriophage T7. Such a finding would implicate the phage ORF product in a related activity. This will also suggest that a

42

new antimicrobial could be derived by a mimetic approach (e.g., peptidomimetic) imitating this function or by a small molecule inhibitor to the bacterial target of the phage ORF, or any steps in the relevant host metabolic pathway, e.g., high throughput screening of small molecule libraries. Selected examples of such similarity between ORFs of Staphyloccus aureus bacteriophage 77 and proteins with inhibitor functions for bacterial hosts are listed in Figure 4 of the provisional application 60/110,992. These include orf9 (similar to bacteriophage P1 kilA function), and orf4 (autolysin of Staphylococcus aureus, amidase enzymatic activity).

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A reason for the biochemical study of individual ORFs for inhibitor function is that their expression or overexpression will block cellular pathways of the host, ultimately leading to arrest and/or inhibition of host metabolism. In addition, such ORFs can alter host metabolism in different ways, including modification of pathogenicity. Therefore, individual ORFs identified above are expressed, preferably overexpressed, in the host and the effect of this expression or overexpression on host metabolism and viability is measured. This approach can be systematically applied to every ORF of the phage, if necessary, and does not rely on the absolute identification of candidate ORFs by bioinformatics. Individual ORFs are resynthesized from the phage genomic DNA, e.g., by the polymerase chain reaction (PCR), preferably using oligonucleotide primers flanking the ORF on either side. These single ORFs are preferably engineered so that they contain appropriate cloning sites at their extremities to allow their introduction into a new bacterial expression plasmid, allowing propagation in a standard bacterial host such as E. coli, but containing the necessary information for plasmid replication in the target microbe such as S. aureus (hereafter referred to as shuttle vector). Shuttle vectors and their use are well known in the art.

Such shuttle vectors preferably also contain regulatory sequences that allow inducible expression of the introduced ORF. As the candidate ORF may encode an inhibitor function that will eliminate the host, it is beneficial that it not be expressed prior to testing for activity. Thus, screening for such sequences when expressed in a constitutive fashion is less likely to be successful when the inhibitor is lethal. In the exemplary inducible system presented in Figure 1A, 1B, 2, and 7, regulatory sequences from the ars operon of S. aureus are used to direct individual ORF expression in S. aureus (or other bacteria in which the ars system is functional). The ars operon encodes a series of proteins which normally mediate the extrusion of arsenite and other trivalent oxyanions from the cells when they are exposed to such toxic substances in their environment. The operon encoding this detoxifying mechanism is normally silent and only induced when arsenite-related compounds are

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present. (Tauriainen, S. et al. (1997) App. Env. Microb., Vol. 63, No. 11, p. 4456-4461.)

Therefore, individual phage ORFs can be expressed in *S. aureus* in an inducible fashion by adding to the culture medium non-toxic arsenite concentrations during the growth of individual *S. aureus* clones expressing such individual phage ORFs. Toxicity of the phage inhibitor ORF for the host is monitored by reduction or arrest of growth under induction conditions, as measured by optical density in liquid culture or after plating the induced cultures on solid medium. Subsequently, interference of the phage ORF with the host biochemical pathways ultimately leading to reduced or arrested host metabolism can be measured by pulse-chase experiments using radiolabeled precursors of either DNA replication, RNA transcription, or protein synthesis. Similar constructs can be made and used for other bacteria using well-known techniques.

Those skilled in the art are familiar with a variety of other inducible systems which can also be used for the controlled expression of phage ORFs, including, for example, lactose (see e.g., Stratagene's LacSwitch<sup>TM</sup>II system; La Jolla, CA) and tetracycline-based systems (see, e.g. Clontech's Tet On/Tet Off<sup>TM</sup> system; Palo Alto, CA). The arsenite-inducible system described is further depicted in Figures 1, 2 and 7.

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The selection or construction of shuttle vectors and the selection and use of inducible systems are well known and thus other shuttle vectors appropriate for other bacteria can be readily provided by those skilled in the art, e.g., for use in other bacterial species.

Standard methodologies for expressing proteins from constructs, and isolating and manipulating those proteins, for example in cross-linking and affinity chromatography studies, may be found in various commonly available and known laboratory manuals. See, e.g., Current Protocols in Protein Science, John Wiley & Sons, Secaucus, N.J., and Maniatis, T. et al. (1989) Molecular Cloning: A Laboratory Manual, Cold Spring Harbor University Press, Cold Spring, N.Y.

It has been found that certain phage or other viruses inhibit host cells, at least in part, by producing an antisense RNA which binds to and inhibits translation from a bacterial RNA sequence. Thus, in the case of potentially inhibitor RNA transcripts encoded by the phage genome, a strong indicator of a possible inhibitory function is provided by the identification of phage sequence which is the identical to or fully complementary (or with only a small percentage of mismatch, e.g., <10%, preferably less than 5%, most preferably less than 3%, to a bacterial sequence. This approach is convenient in the case of bacteria that have been essentially completely sequenced, as the comparison can be performed by computer using public database information.

44

The inhibitory effect of the transcript can be confirmed using expression of the phage sequence in a host bacterium. If needed, such inhibitory can also be tested by transfecting the cells with a vector that will transcribe the phage sequence to form RNA in such manner that the RNA produced will not be translated into a polypeptide. Inhibition under such conditions provides a strong indication that the inhibition is due to the transcript rather than to an encoded polypeptide.

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In an alternative, the expression of an ORF in a host bacterium is found to be inhibitory, but the inhibition is found to be due to an RNA product of the genomic coding region. For antisense inhibition, the sequence of the bacterial target nucleic acid sequence can be identified by inspection of the phage sequence, and the full sequence of the relevant coding region for the bacterial product can be found from a database of the bacterial genomic sequence or can be isolated by standard techniques (e.g., a clone in a genomic library can be isolated which contains the full bacterial ORF, and then sequenced).

In either case, the identification of a target which is inhibited by an RNA transcript produced by a phage provides both the possible inhibition of bacteria naturally containing the same target nucleic acid sequence, as well as the ability to use the target sequence in screening for other types of compounds which will act directly on the target nucleic acid sequence or on a polypeptide product expressed or regulated, at least in part, by the target of the inhibitory phage RNA.

In some cases it will be found that the target of an inhibitory phage RNA or protein has previously been found to be a target of an inhibitory phage RNA or protein has previously been found to be a target for an antibacterial agent. In such cases, the phage inhibitor can still provide useful information if it is found that the phage-encoded product acts at a different site than the previously identified antibacterial agent or inhibitor, i.e., acts at a phage-specific site. For many targets, action at a different site provides highly beneficial characteristics and/or information. For example, an alternate site of inhibitor action can at least partially overcome a resistance mechanism in a bacterium. As an illustration, in many cases, resistance is due, in large part, to altered binding characteristics of the immediate target to the antibacterial agent. The altered binding is due to a structural change which prevents or destabilizes the binding. However, the structural change is frequently quite local, so that compounds which bind at different local sites will b unaffected or affected to a much lesser degree. Indeed, in some cases the local sites will be on a different molecule and so may be completely unaffected by the local structural change creating resistance to the original agent(s). An example of resistance due to altered binding is

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provided by methicillin-resistant *Staphylococcus aureus*, in which the resistance is due to an altered penicillin-binding protein.

In other cases, a new site of action can have improved accessibility as compared to a site acted on by a previously identified agent. This can, for example, assist in allowing effective treatment at lower doses, or in allowing access by a larger range of types of compounds, potentially allowing identification of more potential active agents.

Another advantage is that the structural characteristics of a different site of action will lead to identification and/or development of inhibitors with different structures and different pharmacological parameter. This can allow a greater range of possibilities when selecting an antibacterial agent.

Yet further, different sites often produce different inhibitory characteristics in the target organism. This is commonly the case for multi-domain target proteins. Thus, inhibition targeting an alternate site can produce more efficacious action, e.g., faster killing, slower development of resistance, lower numbers of surviving cells, and different secondary effects (for example, different nutrient utilization).

## Staphylococcus aureus phage 77

As indicated above, the present invention is concerned, in part, with the use of bacteriophage 77 coding sequences and the encoded polypeptides or RNA transcripts to identify bacterial targets for potential new antibacterial agents.

As described, phage 77 ORFs 17, 19, 43, 102, 104, and 182 have been found to have bacteria inhibiting function. Identification of ORFs 17, 19, 43, 102, 104, and 182 and products from the phage which inhibit the host bacterium both provides an inhibitor compound and allows identification of the bacterial target affected by the phage-encoded inhibitor. Such a target is thus identified as a potential target for development of other antibacterial agents or inhibitors and the use of those targets to inhibit those bacteria. As indicated above, even if such a target is not initially identified in a particular bacterium, such a target can still be identified if a homologous target is identified in another bacterium. Usually, but not necessarily, such another bacterium would be a genetically closely related bacterium. Indeed, in some cases, an inhibitor encoded by phage 77 ORF 17, 19, 43, 102, 104, or 182 can also inhibit such a homologous bacterial cellular component.

Possible bacterial target sequences are described herein by reference to sequence source sites. In preferred embodiments, the sequence encoding the target corresponds

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to a *S. aureus* nucleic acid sequence available from numerous sources including *S. aureus* sequences deposited in GenBank, *S. aureus* sequences found in European Patent Application No. 97100110.7 to Human Genome Sciences, Inc. filed January 7, 1997, *S. aureus* sequences available from TIGR at

http://www.tigr.org/tdb/mdb/mdb.html, and S. aureus sequences available from the Oklahoma University S. aureus sequencing project at the following URL:

http://www.genome.ou.edu/staph\_new.html. Such possible targets are particularly applicable to S aureus phages 77, 3A, 96, and 44 AHJD.

The amino acid sequence of a polypeptide target is readily provided by translating the corresponding coding region. For the sake of brevity, the sequences are not reproduced herein. Also, in preferred embodiments, a target sequence corresponds to a S. aureus coding sequence corresponding to a sequence listed in Table 15 herein. The listing in Table 15 describes S. aureus sequences currently listed with GenBank. Again, for the sake of brevity, the sequences are described by reference to the database accession numbers instead of being written out in full herein. In cases where an entry for a coding region is not complete, the complete sequence can be readily obtained by routine methods, e.g., by isolating a clone in a phage host S. aureus genomic library, and sequencing the clone insert to provide the relevant coding region. The boundaries of the coding region can be identified by conventional sequence analysis and/or by expression in a bacterium in which the endogenous copy of the coding region has been inactivated and using subcloning to identify the functional start and stop codons for the coding region.

#### Staphyloccus aureus phage 44 AHJD

The present invention also can utilize the identification of naturally occurring DNA sequence elements within *Staphylococcus aureus* bacteriophage 44AHJD which encode proteins with antimicrobial activity.

Such identification can utilize bioinformatics identification of specific proteins (ORFs) utilized by Staphylococcus aureus bacteriophage 44AHJD during the viral life cycle, resulting in a slowing or arrest of growth of the bacterial host, or in death, of the Staphylococcus aureus host including lysis of the infected bacteria. Thus, some of the bacteriophage 44AHJD DNA sequences encoding these proteins (ORFs) are predicted to encode antimicrobial functions. Information derived from these DNA sequences and translated ORFs can, in turn, be utilized to develop inhibitory compounds by peptidomimetics that can also function as antimicrobials. In addition, the identification of the host bacterial proteins that are targeted and inhibited by the

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antimicrobial bacteriophage ORFs can themselves provide novel targets for drug discovery.

The methodology described above is used to identify and characterize DNA sequences from *Staphylococcus* sp. bacteriophage 44 AHJD that have antimicrobial activity. As described in the Examples, the *Staphylococcus aureus* propagating strain (PS 44A), obtained from the Felix d'Herelle Reference Centre (#HER 1101), was used as a host to propagate its phage 44AHJD, also obtained from the Felix d'Herelle Reference Centre (#HER 101). By sequencing, we found that bacteriophage 44AHJD consists of 16,668 bp (Table 16) predicted to encode 73 ORFs greater than 33 amino acids (Tables 17 & 18). Computational analysis of the predicted protein products of *Staphylococcus aureus* bacteriophage 44AHJD identified homolgs in public sequence databases as listed inTable 19 and 20, along with the accompanying list of related proteins.

From this analysis, it is apparent that 3 genes (ORF 3, 7, and 8) are related to structural proteins found in other bacteriophages. These include genes predicted to encode a tail protein (ORF 3), an upper collar/connector protein of the phage virion (ORF 7), and a lower collar protein (ORF 8). Bioinformatics has also identified one gene whose product is likely involved in phage DNA synthesis. One gene (ORF 1) shows significant homology to DNA polymerases of a number of bacteriophages. bacteria and fungi, and the product of this gene is likely responsible for replicating the genetic material of bacteriophage 44AHJD. ORF 2 encodes a protein with homology to the dinC gene of Bacillus subtilis that encodes a protein involved in teichoic acid biosynthesis. Teichoic acid is a polyphosphate polymer found in some, but not all, Gram positive organisms (and not in Gram negative organisms), where it is attached to the peptidoglycan layer. The phage protein may thus be involved in the synthesis of this material for incorporation into the cell wall, allowing enhanced lysis by the phage lysis enzymes or, as many enzymes can function in "reverse reactions", may be involved in its degradation allowing for penetration of the peptidoglycan and phage genome entry into the cell following adsorption. The similarity between Staphylococcus aureus bacteriophage 44AHJD and E. coli phage T7 indicates that they may share similar mechanisms of replication and growth. Both phages belong to the Pododviridae Family of bacteriophages and are members of the "T7-like" Genus of this Family (Ackermann and DuBow; VIth ICTV Report).

48

Two genes, ORF 9 and 12, were identified with the potential to encode antimicrobial protein products. The homology alignments are shown in Tables 19 and 20. The predicted product of ORF 9 is related to a class of genes which encodes lysozyme-like functions, enzymes which cleave linkages in the mucopolysaccharide cell wall structure of a variety of micro-organisms, including that from the *Staphylococcus aureus* bacteriophage Twort. ORF 12 of *Staphylococcus aureus* bacteriophage 44AHJD shows homology to a set of lysis proteins from several bacteriophages. These lysis proteins are also referred to as holins, and represent phage-encoded lysis functions required for transit of the phage murein hydrolases (lysozyme) to the periplasm, where it can digest the cell wall and thus lyse the bacterium.

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Thus, in particular embodiments, the present invention provides a nucleic acid sequence isolated from Staphylococcus aureus bacteriophage 44AHJD comprising at least a portion of one of the genes described above with antimicrobial activity. For example, ORF 1 encodes a DNA polymerase function. This polymerase may utilize host-derived accessory proteins for its activity when replicating the phage template, sequestering such proteins from use by the bacterial polymerase, resulting in inhibition of DNA replication, cell division, and cell growth. Alternatively, ORF 9 directly encodes a polypeptide with antimicrobial activity. ORF 9 is predicted to encode an amidase, a protein known to act as a cell wall degrading enzyme. ORF 12 likely encodes a holin function required for transit of the phage amidase (gene 9) product) to the periplasm. When this type of gene product from Bacillus phage phi 29 (gene 14), was cloned in Escherichia coli, cell death ensued (Steiner et al., 1993). Thus, production of proteins from Bacillus phage phi 29 gene 14 in E. coli resulted in cell death, whereas production of protein from Bacillus phage phi 29 gene 14 concomitantly with the phi 29 lysozyme or unrelated murein-degrading enzymes led to lysis, suggesting that membrane-bound protein 14 induces a nonspecific lesion in the cytoplasmic membrane (Steiner et al., 1993).

The present invention also provides the use of the *Staphylococcus* bacteriophage 44 AHJD antimicrobial ORFs or ORF products as pharmacological agents, either wholly or in part and derivatives, as well as the use of corresponding peptidomimetics, developed from amino acid or nucleotide sequence knowledge derived from *Staphylococcus* bacteriophage 44 AHJD killer ORFs.

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Enterococcus phage 182

Bacteriophage 182 was obtained from the Felix D'Herelle phage collection (Ste. Foy, Quebec) and infects *Enterococcus sp.* Group D. The genome of *Enterococcus* bacteriophage 182 consists of 17,833 bp (Table 21) and is predicted to encode 80 ORFs greater than 33 amino acids (Tables 22 and 23). Computational analysis of the predicted protein products of *Enterococcus* bacteriophage 182 was performed in order to identify protein products related to those deposited in public databases. Bacteriophage 182 protein products which detected sequences with significant sequence similarity in public databases are listed in Table 24 and 26, along with the accompanying list of related proteins.

From this analysis, it is apparent that 5 genes (ORF 001, 004, 007, 009, and 011) are related to structural proteins of several *Bacillus* phages – *Bacillus* bacteriophage PZA, phi-29, and B103. These include genes predicted to encode a tail protein (ORF 001), a head protein (ORF 004), and upper collar protein (ORF 007), a lower collar protein (ORF 009), and a pre-neck appendage protein (ORF 011). Two gene products are predicted to encode genes which direct phage morphogenesis – these are ORF 005 and 019.

Bioinformatics has also identified three genes whose products are likely involved in phage DNA synthesis. One gene, ORF 002 shows significant homology to DNA polymerases of a number of bacteriophages, and the product of this gene is likely responsible for replicating the genetic material of bacteriophage 182. ORF 006 encodes a protein with homology to the encapsidation proteins of several other bacteriophages, including *Bacillus* phage phi-29 (P11014), PZA (P07541), and B103 (X99260) and *Streptococcus* phage CP-1 (Z47794). These gene products catalyze the *in vivo* and *in vitro* genome-encapsidation reaction (Garvey et al., 1985). Proteins involved in genome packaging have been shown to have additional activities that affect biochemical reactions in other phages and their hosts. For example, the coat protein of the RNA bacteriophage MS2 interacts with viral RNA to translationally repress replicase synthesis (Pickett and Peabody, 1993). This protein-RNA interaction also plays a role in genome encapsidation, enveloping a single copy of the viral genome in a protein shell composed of many molecules of coat protein. In addition, the bacteriophage λ terminase enzyme can be lethal to *E. coli* when expressed,

suggesting cleavage of packaging sites in the bacterial chromosome. Also present within bacteriophage 182 is a gene, ORF 010, that encodes a protein that is related to the terminal proteins of *Bacillus* phage Nf (P06812), *Bacillus* phage GA-1 (X96987) and *Bacillus* phage B103 (X99260). DNA terminal proteins are linked to the 5' ends of both strands of the genome and are essential for DNA replication playing a role in initial priming of DNA replication. The similarity between *Enterococcus* bacteriophage 182 and Bacillus phages phi-29, PZA, and B103 indicates that they may share similar mechanisms of replication and growth. Protein-primed DNA replication is a well described phenomenon, and in the phi-29-like phages, the ends of the DNA serve as origins and termini of replication (Gutiérrez et al., 1986; Yoshikawa et al., 1985).

There is also a gene (ORF 015) that encodes a protein showing homology to an early protein product of *Bacillus* bacteriophage PZA and the single-strand nucleic acid binding protein of bacteriophage B103.

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Two genes, ORF 008 and 014, were identified with the potential to encode anti-microbial protein products. The homology alignments are shown in Tables 24 & 26 and biochemical features of the predicted polypeptides shown in Table 25. The predicted product of ORF 008 is related to a class of genes which encodes lysozyme-like functions, enzymes which cleave linkages in the mucopolysaccharide cell wall structure of a variety of micro-organisms. ORF 014 of *Enterococcus* 182 shows homology to a set of lysis proteins from *Bacillus* bacteriophage phi-29, PZA, and B103. These lysis proteins are also referred to as holins and represent phage encoded lysis functions required for transit of the phage murein hydrolases (lysozyme) to the periplasm, where it can digest the outer cell wall and thus lyse the bacterium.

Thus, the present invention provides a nucleic acid sequence obtained from *Enterococcus* bacteriophage 182 comprising at least a portion of a phage 182 ORF, preferably an inhibitory ORF, and more preferably at least a portion of one of the genes described above with anti-microbial activity. For example, ORF 002 encodes a DNA polymerase function. This polymerase may utilize host-derived accessory proteins for its activity when replicating the phage template, sequestering such proteins from use by the bacterial polymerase, resulting in inhibition of DNA replication, cell division, and cell growth. Alternatively, ORFs 008 or 014 directly encode polypeptides with anti-microbial activity. ORF 008 is predicted to encode an

51

autolytic lysozyme, a protein known to have anti-microbial activity (Martin et al., 1998). ORF 014 likely encodes a holin function required for transit of the phage murein hydrolases to the periplasm. When the related product from Bacillus phage phi 29 (gene 14), was cloned in Escherichia coli, cell death ensued (Steiner et al., 1993). Thus, production of proteins from Bacillus phage phi 29 gene 14 in E. coli resulted in cell death, whereas production of protein from Bacillus phage phi 29 gene 14 concomitantly with the phi 29 lysozyme or unrelated murein-degrading enzymes led to lysis, suggesting that membrane-bound protein 14 induces a nonspecific lesion in the cytoplasmic membrane (Steiner et al., 1993).

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The present invention also provides the use of the Enterococcus bacteriophage 182 anti-microbial ORFs as pharmacological agents, either wholly or in part and derivatives, as well as the use of corresponding peptidomimetics, developed from amino acid or nucleotide sequence knowledge derived from Enterococcus bacteriophage 182 killer ORFs. This can be done where the structure of the peptidomimetic compound corresponds to the structure of the active portion of a product of an ORF. In this analysis, the peptide backbone is transformed into a carbon based hydrophobic structure that can retain cytostatic or cytocidal activity for the bacterium. This is done by standard medicinal chemistry methods, measuring growth inhibition of the various molecules in liquid cultures or on solid medium. These mimetics also represent lead compounds for the development of novel antibiotics. In this context, "corresponds" means that the peptidomimetic compound structure has sufficient similarities to the structure of the active portion of a product of one of the Enterococcus ORFs listed, that the peptidomimetic will interact with the same molecule as the product of the ORF, and preferably will elicit at least one cellular response in common which relates to the inhibition of the cell by the phage protein.

To validate the identity of an ORF as a killer ORF, it is preferably expressed in the host or other test bacterial organism and the effect of this expression on bacterial growth and replication is assessed. Therefore, all individual ORFs identified herein, e.g., those identified above, can be expressed, preferably overexpressed, in a suitable host bacterium e.g., a host *Enterococcus* and the effect of this expression or overexpression on host metabolism and viability can be measured.

Individual ORFs can be resynthesized from the phage genomic DNA by the polymerase chain reaction (PCR) using oligonucleotide primers flanking the ORF on

52

either side. Those skilled in the art are familiar with the design and synthesis of appropriate primer sequences. These single ORFs are preferably engineered so that they contain appropriate cloning sites at their extremities to allow their introduction into a new bacterial expression plasmid, allowing propagation in a standard bacterial host such as *E. coli*, but containing the necessary information for plasmid replication in the target microbe, *Enterococcus* sp. (hereafter referred to as a shuttle vector).

This shuttle vector also preferably contains regulatory sequences that allow inducible expression of the introduced ORF. As the candidate ORF may encode a killer function that will eliminate the host, it is highly advantageous that it not be expressed (or at least not expressed at a substantial level) prior to testing for activity; thus screening for such sequences in a constitutive fashion is less likely to be successful (lethality). In an example presented in Fig. 7, regulatory sequences from the ars operon are used to direct individual ORF expression in Enterococcus. The ars operon encodes a series of proteins which normally mediate the extrusion of arsenite and several other trivalent oxyanions from the cells when they are exposed to such toxic substances in their environment. The operon encoding this detoxifying mechanism is normally silent and only induced when arsenite-related compounds are present.

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Therefore, individual phage ORFs can be expressed in *Enterococcus* or other suitable host in an inducible fashion by adding to the culture medium non-toxic arsenite concentrations during the growth of individual *Enterococcus* (or other host cells) clones expressing such individual phage ORFs. Toxicity of the phage killer ORF for the host is monitored by reduction or arrest of growth under induction conditions, as measured by optical density in liquid culture or after plating the induced cultures on solid medium. Subsequently, interference of the phage ORF with the host biochemical pathways ultimately leading to reducing or arresting host metabolism can be measured by pulse chase experiments using radiolabeled precursors of either DNA replication, RNA transcription, or protein synthesis.

Of course, other inducible regulatory sequences (e.g., promoters, operators, etc.) may be used (e.g., systems using positive induction of expression or systems using release of repression). A variety of such systems are known to those-skilled in the art and can be utilized in the present invention.

Nucleic acid sequences of the present invention can be isolated using a method similar to those described herein or other methods known to those skilled in the art. In addition, such nucleic acid sequences can be chemically synthesized by well-known methods. Having the phage 182 ORFs, e.g., anti-bacterial ORFs of the present invention, portions thereof, or oligonucleotides derived therefrom as described, other anti-microbial sequences from other bacteriophage sources can be identified and isolated using methods described here or other methods, including methods utilizing nucleic acid hybridization and/or computer-based sequence alignment methods.

The invention also provides bacteriophage anti-microbial DNA segments from other phages based on nucleic acids and sequences hybridizing to the presently identified inhibitory ORF under high stringency conditions or sequences which are highly homologous. The bacteriophage anti-microbial DNA segment from bacteriophage 182 can be used to identify a related segment from another unrelated phage based on stringent conditions of hybridization or on being a homolog based on nucleic acid and/or amino acid sequence comparisons. As with the phage 182 inhibitory sequences, such homologous coding sequences and products can be used as antimicrobials, to construct active portions or derivatives, to construct peptidomimetics, and to identify bacterial targets.

Enterococcus sequences are listed in Table 27 by accession number, providing identification of possible targets of Enterococcus phage inhibitory ORF products, e.g., from phage 182.

#### Streptococcus pneumoniae

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As indicated in the Summary above, the present invention is concerned with the use of *Streptococcus* sp. bacteriophage Dp-1 coding sequences and the encoded polypeptides or RNA transcripts to identify bacterial targets for potential new antibacterial agents.

Streptococcus pneumoniae is an important cause of community-acquired pneumonia and a major cause of otitis media, sinusitis, and meningitis in children and adults. In Spain and other Mediterranean countries, the majority of S. pneumoniae are relatively resistant to penicillin (Klugman, 1990; Fenoll et al., 1991; Jorgensen et al., 1990). These strains also have decreased susceptibility to broad-spectrum cephaloporins, which are frequently used in the empiric treatment of meningitis and

54

other serious invasive bacterial infections. High-level resistance of pneumococci has been encountered in Hungary where 70% of children who were colonized with *S. pneumoniae* carried penicillin resistant strains that were also resistant to tetracycline, erythromycin, trimethoprim/sulfamethoxazole, and 30% resistant to chloramphenicol (Neu, 1992). The resistance of pneumococci to macrolides such as erythromycin averages 20-25% in France, ~20% in Japan, and <10% in Spain (Neu, 1992).

The antimicrobial susceptibilities and distribution of serotypes of the 42 isolates of *S. pneumoniae* in southern Taiwan from invasive infections have been recently determined (Hseuh et al., 1996). Resistance rates among these isolates were: erythromycin, 61.9%; clindamycin, 47.6%; chloramphenicol, 19%; and tetracycline, 73.8%. Resistance to three or more classes of antibiotics was found in 33.3% of the isolates. Bacteremic pneumonia and primary bacteremia accounted for 64.3% of the infections and mortality was 42.6%. Given the severity of these infections despite adequate antibiotic therapy, there is clearly a need for introduction of new therapeutic options to prevent mortality due to invasive *S. pneumoniae* infections.

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Pneumococcal phages belong to four families and they present a great variety in morphology, including lytic and temperate phages (for a review, see Garcia et al., 1997). Examples of lytic phages are Cp-1 and Dp-1, whereas examples of temperate phages are HB-3, EJ-1, and HB-746. The complete nucleotide sequence and functional organization of Cp-1 has been reported (Martin et al., 1996). Cp-1 has a 19,345 bp double-stranded DNA genome, with a terminal protein covalently linked to its 5' ends, that replicates by a protein primed mechanism. The phage contains 29 ORFs, 23 on one strand and 6 on the opposite. When these predicted proteins were compared to sequences compiled in GenBank EMBL databases, to ORFs showed significant similarity to proteins of bacteriophage 29 that infects B. subtilis (Martin et al., 1996). The similar proteins corresponded to those involved in DNA replication (terminal protein and DNA polymerase), structural and morphogenic proteins (major head, collar, connector, tail, and encapsidation proteins), and proteins involved in lysis function (holin and lysozyme). In its strategy of lysis, the holin gene product inserts itself into the cell membrane, allowing access of the lysozyme to the peptidoglycan. Expression of the Cp-1 holin protein in E. coli results in cell death after 2-hours of induction, but did not lead to lysis (Garcia et al., 1997). Cells harboring a plasmid construction with holin and lysozyme genes together did lyse after induction and the

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viability loss was similar to that of the culture expressing holin alone. Cloning of these lytic genes in *S. pneumoniae* showed that both genes had the same effect as in *E. coli*. That is, holin itself did not lyse the culture but the viability loss was noticeable, whereas both holin and lysozyme together were capable of lysing M31, an amidase deleted mutant (Garcia et al., 1997).

Recently, a small portion (~4 kbp) of a second *S. pneumoniae* phage, Dp-1, has been sequenced (Sheehan et al., 1997). This portion contains the genes coding for the lytic system (Sheehan et al., 1997) and shows a modular organization similar to that described for Cp-1. However, in this case, a single chimeric protein appears to be made in which the N-terminal domain is highly similar to that of the murein hydrolase coded by a gene found in the phage BK5-T that infects *Lactococcus lactis*, and the C-terminal domain is homologous to holins. Thus, both functions appear to have been combined in a novel chimeric protein.

Bacteriophage Dp-1 was obtained from Dr. P. Garcia (Departamento de Microbiologia Molecular, Centro de Departamento de Investigaciones Biologicas, Consejo Superior de Investigaciones Cientificas, Velazquez, Madrid, Spain). We found that Dp-1 has a double-stranded DNA genome of 56,506 bp, predicted to encode 85 ORFs greater than 33 amino acids and with upstream Shine-Dalgarno motifs for translation initiation (Tables 28 & 30, and Fig. 6). Computational analysis of the predicted protein products of *Streptococcus* bacteriophage Dp-1 protein products, which detected homologs in public databases, are listed inTable 31, along with the accompanying list of related proteins.

From this analysis, it is apparent that several predicted genes of Dp-1 encode polypeptides that are related to structural proteins. ORFs 001, 002, 004, and 030 are predicted to encode tail proteins, minor structural proteins, and minor capsid proteins (Table 31). We also note the identification of several gene products that are likely involved in DNA synthesis. These include ORF 3 which encodes DNA polymerase, ORF 8 which encodes a SWI/SNF helicase-related protein, ORF 10 encodes a protein showing homology to recA, and ORF 13 encodes a dnaZX-like ORF.

In *E. coli*, RapA encodes an RNA polymerase (RNAP)-associated protein with ATPase activity and which is a homolog of the eukaryotic SWI/SNF family, a set of proteins whose members are involved are involved in transcription activation, nucleosome remodeling, and DNA repair. RapA forms a stable complex with RNAP,

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as if it were a subunit of RNAP and it is possible that the ORF 8 product behaves similarly or in a dominant-negative fashion to inhibit the activity of RapA. Mutation of the essential *E. coli* dnaZX results in a block in DNA chain elongation during replication (Maki et al., 1988). The dnaZX gene has only one open reading frame for a 71-kDa polypeptide from which the two distinct DNA polymerase III holoenzyme subunits, tau (71 kDa) and gamma (47 kDa), are produced. The tau subunit is the precursor of the gamma subunit, and the gamma subunit is produced by a -1 frameshift causing early termination of translation (Tsuchihashi et al., 1990). These proteins show single-strand DNA binding properties that is ATPase (and dATPase) dependent and are thought to increasing the processivity of the core DNA polymerase enzyme (Lee et al., 1987).

There are several Dp-1 ORFs which encode proteins predicted to play a role in cellular metabolic pathways. These include polypeptides involved in coenzyme PQQ synthesis (ORFs 20, 29, 38). Pyrrolo-quinoline quinone (PQQ) is the non-covalently bound prosthetic group of many quinoproteins catalysing reactions in the periplasm of Gram-negative bacteria. Most of these involve the oxidation of alcohols or aldose sugars. Interestingly, ORFs 20, 29, and 30 also show homology to the exoenzyme S regulon (Frank, 1997). Proteins encoded by the *P. aeruginosa* exoenzyme S regulon may be involved in a contact-mediated translocation mechanism to transfer anti-host factors directly into eukaryotic cells disrupting eukaryotic signal transduction through ADP-ribosylation (Frank, 1997).

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There is also a protein with similarity to GTP cyclohydrolase I (ORF 21) and ORF 41 which shows homology to dUTPase (Table 31). GTP cyclohydrolase I is an enzyme that catalyzes the first reaction in the pathway for the biosynthesis of the pteridine, a cofactor of the monooxygenases of the aromatic amino acids. Disruption of the homologous gene in *Saccharomyces cerevisiae* leads to a recessive conditional lethality due to folinic acid auxotrophy, that can be complemented with the mammalian or bacterial GTP cyclohydrolase I enzymes (Nardese et al., 1996; Mancini et al., 1999).

ORF 16 shows high homology to autolysin. This region of the phage sequence was previously reported (Sheehan et al., 1997) and encompasses ~ 4 kbp of our sequence. The sequence published by (Sheehan et al., 1997) is shown in Table 32.

Thus, the present invention provides a nucleic acid sequence obtained from *Streptococcus* bacteriophage Dp-1 comprising at least a portion of a phage Dp-1 ORF, preferably an inhibitory ORF, and more preferably at least a portion of one of the genes described above with anti-microbial activity. For example, ORF 013 encodes a

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protein with homology to the gamma subunit of DNA polymerase (dnaX gene). This protein may act in a dominant-negative fashion to sequester the host DNA polymerase for its own replication, thus inhibiting host DNA replication. The dnaX gene product is essential for *E. coli* replication (Kodaira et al., 1983).

In certain preferred embodiments of the present invention, the bacterial target of a bacteriophage inhibitor ORF product, e.g., an inhibitory protein or polypeptide, is encoded by a *Streptococcus* nucleic acid coding sequence from a host bacterium for bacteriophage Dp-1. As above, possible target sequences are described herein by reference to sequence source sites. The sequence encoding the target preferably corresponds to a *Streptococcus* nucleic acid sequence available from The Institute for Genomic Research (TIGR), or available from GenBank or other public database. The TIGR *Streptococcus* sequences are publicly available at The Institute for Genomics Research at URL: <a href="http://www.tigr.org">http://www.tigr.org</a>

The amino acid sequence of a polypeptide target is readily provided by translating the corresponding coding region. For the sake of brevity, the sequences are not reproduced herein. Also, in preferred embodiments, a target sequence corresponds to a Streptococcus pneumoniae coding sequences corresponding to a sequence listed in Table 33 herein. Sequences for other Streptococcal species are also available from TIGR and./or from GenBank. The listing in Table 33 describes Streptococcus sequences currently deposited in GenBank. Again, for the sake of brevity, the sequences are described by reference to the GenBank entries instead of being written out in full herein. In cases where the TIGR or GenBank entry for a coding region is not complete, the complete sequence can be readily obtained by routine methods, e.g., by isolating a clone in a phage Dp-1 host Streptococcus sp. genomic library, and sequencing the clone insert to provide the relevant coding region. The boundaries of the coding region can be identified by conventional sequence analysis and/or by expression in a bacterium in which the endogenous copy of the coding region has been inactivated and using subcloning to identify the functional start and stop codons for the coding region.

In the various aspects of this invention involving Dp-1 sequences, preferably the sequence is preferably not contained in the sequence described in Sheehan et al., 1997 (Table 32).

## Validating Identified Inhibitory Phage ORFs

A fifth step involves validating the identified phage inhibitor ORF by independent methods, and delineating further possible smaller segments of the ORFs

that have inhibitory activity. Several methods exist to validate the role of the identified ORF as an inhibitor ORF.

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One example utilizes the creation of a mutant variant of the phage ORF in which the candidate ORF carries a partial or complete loss-of-function mutation that is measurable as compared with the non-mutant ORF. Comparison of the effects of expression of the loss of function mutant with the normal ORF provides confirmation of the identification of an inhibitor ORF where the loss-of-function mutant provides a measurably lower level of inhibition, preferably no inhibition. The loss of function may be conditional, e.g., temperature sensitive.

Once validation of the inhibitor ORF is achieved, a bi-directional deletion analysis can be carried out using the same experimental system to identify the minimal polypeptide segment that has inhibitor activity. This may be carried out by a variety of means, e.g., by exonuclease or PCR methodologies, and is used to determine if a relatively small segment of the ORF (i.e., the product of the ORF) still possesses inhibitory activity when isolated away from its native sequence. If so, a portion of the ORF encoding this "active portion" can be used as a template for the synthesis of novel anti-microbial agents and further allowing derivation of the peptide sequence, e.g., using modified peptides and/or peptidomimetics.

In creation of certain peptidomimetics, the peptide backbone is transformed into a carbon-based hydrophobic structure that can retain inhibitor activity against the bacterium. This is done by standard medicinal chemistry methods, typically monitored by measuring growth inhibition of the various molecules in liquid cultures or on solid medium. These mimetics can also represent lead compounds for the development of novel antibiotics.

Recently, a major effort has been undertaken by the pharmaceutical industry and their biotechnology partners for the sequencing of bacterial pathogen genomes. The rationale is that the systematic sequencing of the genome will identify all of the bacterial proteins and therefore this proteome will be the target for designing novel inhibitor antibiotics. Although systematic, this approach has several major problems. The first is that analysis of primary amino acid sequences of bacterial proteins does not immediately reveal which protein will be essential for viability of the bacterium, and target validation is thus a major issue. The second problem is one of redundancy, as several biochemical pathways are either structurally duplicated in bacteria (different isoforms of the same enzyme), or functionally duplicated by the presence of salvage pathways in the event of a metabolic block in one pathway (different nutritional conditions). The third is that even a valid target may not be structurally or

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functionally amenable to inhibition by small molecules because of inaccessibility (sequestration of target).

Therefore, there is considerable interest within the pharmaceutical and biotechnology industry in identifying key targets for drug discovery amongst the mass of novel targets generated by large-scale genomic sequencing projects.

On the other hand, and underscoring the instant invention, the phages herein described have, over millions of years, evolved specific mechanisms to target such key biochemical pathways and proteins. In the few cases where inhibition by phages has been elucidated (e.g., see ref. 3), such bacterial targets are invariably rate-limiting in their respective biochemical pathways, are not redundant, and/or are readily accessible for inhibition by the phage (or by another inhibitory compound). Therefore, the sixth step of this invention involves identifying the host biochemical pathways and proteins that are targeted by the phage inhibitory mechanisms.

# 15 <u>Identifying, Validating, and Characterizing Bacterial Host Target Proteins and</u> <u>Affected Pathways</u>

A rationale for this step is that the inhibitor ORF product from the phage physically interacts with and/or modifies certain microbial host components to block their function. Exemplary approaches which can be used to identify the host bacterial pathways and proteins that interact with, and preferably also are inhibited by, phage ORF product(s) are described below.

One approach is a genetic screen to determine physiological protein:protein interaction, for example, using a yeast two hybrid system. In this assay, the phage ORF is fused to the carboxyl terminus of the yeast Gal4 activation domain II (amino acids 768-881) to create a bait vector. A cDNA library of cloned S. aureus sequences which have been engineered into a plasmid where the S. aureus sequences are fused to the DNA binding domain of Gal4 is also generated. These plasmids are introduced alone, or in combination, into yeast strain Y190 - previously engineered with chromosomally integrated copies of the E. coli lacZ and the selectable HIS3 genes, both under Gal4 regulation (Durfee, T., Becherer, K., Chen, P.-L., Yeh, S.-H., Yang, Y., Kilburn, A.E., Lee, W.-H., and Elledge, S.J. (1993). Genes & Dev. 7, 555-569). If the two proteins expressed in yeast interact, the resulting complex will activate transcription from promoters containing Gal4 binding sites. A lacZ and His3 gene, each driven by a promoter containing Gal4 binding sites, have been integrated into the. genome of the host yeast system used for measuring protein-protein interactions. Such a system provides a physiological environment in which to detect potential protein interactions. This system has been extensively used to identify novel protein-protein

interaction partners and to map the sites required for interaction (for example, to identify interacting partners of translation factors (Qiu, H., Garcia-Barrio, M.T., and Hinnebusch, A.G. (1998). *Mol & Cell Biology* 18, 2697-2711), transcription factors (Katagiri, T., Saito, H., Shinohara, A., Ogawa, H., Kamada, N., Nakamura, Y., and Miki, Y. (1998). Genes, *Chromosomes & Cancer* 21, 217-222), and proteins involved in signal transduction (Endo, T.A., Masuhara, M., Yokouchi, M., Suzuki, R., Sakamoto, H., Mitsui, K., Matsumoto, A., Tanimura, S., Ohtsubo, M., Misawa, H., Miyazaki, T., Leonor N., Taniguchi, T., Fujita, T., Kanakura, Y., Komiya, S., and Yoshimura, A. *Nature*. 387, 921-924). This approach has also been used in many published reports to identify interaction between mammalian viral and mammalian cell proteins.

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For example, the non-structural protein NS1 of parvovirus is essential for viral DNA amplification and gene expression and is also the major cytopathic effector of these viruses. A yeast two-hybrid screen with NS1 identified a novel cellular protein of unknown function that interacts with NS-1, called SGT, for small glutamine-rich tetratricopeptide repeat (TPR)-containing protein (Cziepluch C. Kordes E. Poirey R. Grewenig A. Rommelaere, J, and Jauniaux JC. (1998) *J Virol.* 72, 4149-4156). In another screen, the adenovirus E3 protein was recently shown to interact with a novel tumor necrosis factor alpha-inducible protein and to modulate some of the activities of E3 (Li Y. Kang J. and Horwitz M.S. (1998). *Mol & Cell Biol.* 18, 1601-1610). In yet another recent screen, the herpes simplex virus 1 alpha regulatory protein ICP0 was found to interact with (and stabilize) the cell cycle regulator cyclin D3 (Kawaguchi Y. Van Sant C. and Roizman B. (1997). *J Virol.* 71,7328-7336).

Another two-hybrid system for identifying protein:protein interactions is 25 commercially available from STRATEGENE™ as the CYTO-TRAP™ system (Chang et al., Strategies Newsletter 11(3), 65-68 (1998)(from Stratagene)). The system is a yeast-based method for detecting protein:protein interactions in vivo, using activation of the Ras signal transduction cascade by localizing a signal pathway component, human Sos (hSos), to its activation site in the yeast plasma membrane. 30 The system uses a temperature-sensitive Saccharomyces cerevisiae mutant, strain cdc25H, which contains a point mutation at amino acid residue 1328 of the cdc25 gene. This gene encodes a guanyl nucleotide exchange factor which binds and activates Ras, leading to cell growth. The mutation in the cdc25 gene prevents host growth at 37°C, but at a permissive temperature of 25°C, growth is normal. The 35 system utilizes the ability of (hSos) to complement the cdc25 defect and activate the yeast Ras signaling pathway. Once (hSos) is expressed and localized to the plasma membrane, the cdc25H yeast strain grows at 37°C. Localizing hSos to the plasma

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membrane occurs through a protein:protein interaction. A protein of interest, or bait, is expressed as a fusion protein with hSos. The library, or target proteins are expressed with the myristylation membrane-localization signal. The yeast cells are then incubated under restrictive conditions (37°C). If the bait and the target protein interact, the hSos protein is recruited to the membrane, activating the Ras signaling pathway and allowing the cdc25H yeast strain to grow at the restrictive temperature.

The protein targets of phage inhibitory ORFs can also be identified using bacterial genetic screens. One approach involves the overexpression of a phage inhibitory protein in mutagenized bacterial host species, followed by plating the cells and searching for colonies that can survive the antimicrobial activity of the inhibitory ORF. These colonies are then grown, their DNA extracted, and cloned into an expression vector that contains a replicon of a different incompatibility group from the plasmid expressing the original ORF. This library is then introduced into a wild-type host bacterium in conjunction with an expression vector driving synthesis of the phage ORF, followed by selection for surviving bacteria. Thus, bacterial DNA fragments from the survivors presumably contain a DNA fragment from the original mutagenized host bacterial genome that can protect the cell from the antimicrobial activity of the inhibitory phage ORF. This fragment can be sequenced and compared with that of the bacterial host to determine in which gene the mutation lies. This approach enables one to determine the targets and pathways that are affected by the killing function.

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A second approach is based on identifying protein:protein interactions between the phage ORF product and bacterial S. aureus, e.g., proteins using a biochemical approach based, for example, on affinity chromatography. This approach has been used, for example, to identify interactions between lambda phage proteins and proteins from their E. coli host (Sopta, M., Carthew, R.W., and Greenblatt, J. (1985) J. Biol. Chem. 260, 10353-10369). The phage ORF is fused to a peptide tag (e.g. glutathione-S-transferase ("GST"), 6xHIS, ("HIS") and/or calmodulin binding protein ("CPB")) within a commercially available plasmid vector that directs high level expression on induction of a suitably responsive promoter driving the fusion's expression. The translated fusion protein is expressed in E. coli, purified, and immobilized on a solid phase matrix via, for example the tag. Total cell extracts from the host bacterium, e.g., S. aureus, are then passed through the affinity matrix containing the immobilized phage ORF fusion protein; host proteins retained on the column are then eluted under different conditions of ionic strength, pH, detergents etc., and characterized by gel electrophoresis and other techniques. Appropriate controls are run to guard against nonspecific binding to the resin. Target proteins thus

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recovered should be enriched for the phage protein/peptide of interest and are subsequently electrophoretically or otherwise separated, purified, sequenced, or biochemically analyzed. Usually sequencing entails individual digestion of the proteins to completion with a protease (e.g.-trypsin), followed by molecular mass and amino acid composition and sequence determination using, for example, mass spectrometry, e.g., by MALDI-TOF technology (Qin, J., Fenyo, D., Zhao, Y., Hall, W.W., Chao, D.M., Wilson, C.J., Young, R.A. and Chait, B.T. (1997). Anal. Chem. 69, 3995-4001).

The sequence of the individual peptides from a single protein are then analyzed by the bioinformatics approach described above to identify the *S. aureus* protein interacting with the phage ORF. This analysis is performed by a computer search of the *S. aureus* genome for an identified sequence. Alternatively, all tryptic peptide fragments of the *S. aureus* genome can be predicted by computer software, and the molecular mass of such fragments compared to the molecular mass of the peptides obtained from each interacting protein eluted from the affinity matrix. The responsible gene sequence can be obtained, for example by using synthetic degenerate nucleic acid sequences to pull out the corresponding homologous bacterial sequence. Alternatively, antibodies can be generated against the peptide and used to isolate nascent peptide/mRNA transcript complexes, from which the mRNA can be reverse transcribed, cloned, and further characterized using the procedures discussed herein.

A variety of other binding assay methods are known in the art and can be used to identify interactions between phage proteins and bacterial proteins or other bacterial cell components. Such methods that allow or provide identification of the bacterial component can be used in this invention for identifying putative targets.

Validation of the interaction between the phage ORF product and the bacterial proteins or other components can be obtained by a second independent assay (e.g., co-immunoprecipitation or protein-protein crosslinking experiments (Qiu, H., Garcia-Barrio, M.T., and Hinnebusch, A.G. (1998). Mol & Cell Biology 18, 2697-2711; Brown, S. and Blumenthal, T. (1976). Proc. Natl. Acad. Sci. USA 73, 1131-1135)).

Finally, the essential nature of the identified bacterial proteins is preferably determined genetically by creating a constitutive or inducible partial or complete loss-of-function mutation in the gene encoding the identified interacting bacterial protein. This mutant is then tested for bacterial survival and replication.

The protein target of the phage inhibitor function can also be identified using a genetic approach. Two exemplary approaches will be delineated here. The first approach involves the overexpression of a predetermined phage inhibitor protein in mutagenized host bacteria, e.g., S. aureus, followed by plating the cells and searching

for colonies that can survive the inhibitor. These colonies will then be grown, their DNA extracted and cloned into an expression vector that contains a replicon of a different incompatibility group, and preferably having a different selectible marker than the plasmid expressing the phage inhibitor. Thus, host DNA fragments from the mutant that can protect the cell from phage ORF inhibition can be sequenced and compared with that of the bacterial host to determine in which gene the mutation lies. This approach allows rapid determination of the targets and pathways that are affected by the inhibitor.

Alternatively, the bacterial targets can be determined in the absence of selecting for mutations using an approach known as "multicopy suppression". In this approach, the DNA from the wild type host is cloned into an expression vector that can coexist, as previously described, with one containing a predetermined phage inhibitor. Those plasmids that contain host DNA fragments and genes that protect the host from the phage inhibitor can then be isolated and sequenced to identify putative targets and pathways in the host bacteria.

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Regardless of the specific mode of identification, screening assays may additionally utilize gene fusions to specific "reporter genes" to identify a bacterial gene(s) whose expression is affected when the host target pathway is affected by the phage inhibitor. Such gene fusions can be used to search a number of small molecule compounds for inhibitors that may affect this pathway and thus cause cell inhibition. This approach will allow the screening of a large number of molecules on petri dishes or 96-well format by monitoring for a simple color change in the bacterial colonies. In this manner, we can validate host targets and classes of compounds for further study and clinical development. These inhibitors also represent lead compounds for the development of other antibiotics.

Bioinformatics and comparative genomics are preferably then applied to the identified bacterial gene products to predict biochemical function. The biochemical activity of the protein can be verified *in vitro* in cell free assays or *in vivo* in intact cells. *In vitro* biochemical assays utilizing cell-free extracts or purified protein are established as a basis for the screening and development of inhibitors.

These inhibitors, preferably small molecule inhibitors, may comprise peptides, antibodies, products from natural sources such as fungal or plant extracts or small molecule organic compounds. In general, small molecule organic compounds are preferred. These compounds may, for example, be identified within large compound libraries, including combinatorial libraries. For example, a plurality of compounds, preferably a large number of compounds can be screened to determine whether any of the compounds binds or otherwise disrupts or inhibits the identified bacterial target.

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Compounds identified as having any of these activities can then be evaluated further in cell culture and/or animal model systems to determine the pharmacological properties of the compound, including the specific anti-microbial ability of the compound.

For mixtures of natural products, including crude preparations, once a preparation or fraction of a preparation is shown the have an anti-microbial activity, the active substance can be isolated and identified using techniques well known in the art, if the compound is not already available in a purified form.

Identified compounds possessing anti-microbial activity and similar compounds having structural similarity can be further evaluated and, if necessary, derivatized according to synthesis and/or modification methods available in the art selected as appropriate for the particular starting molecule.

#### Derivatization of identified anti-microbials

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In cases where the identified anti-microbials above might represent peptidal compunds, the *in vivo* effectiveness of such compounds may be advantageously enhanced by chemical modification using the natural polypeptide as a starting point and incorporating changes that provide advantages for use, for example, increased stability to proteolytic degradation, reduced antigenicity, improved tissue penetration, and/or improved delivery characteristics.

In addition to active modifications and derivative creations, it can also be useful to provide inactive modifications or derivatives for use as negative controls or introduction of immunologic tolerance. For example, a biologically inactive derivative which has essentially the same epitopes as the corresponding natural antimicrobial can be used to induce immunological tolerance in a patient being treated. The induction of tolerance can then allow uninterrupted treatment with the active anti-microbial to continue for a significantly longer period of time.

Modified anti-microbial polypeptides and derivatives can be produced using a number of different types of modifications to the amino acid chain. Many such methods are known to those skilled in the art. The changes can include, for example, reduction of the size of the molecule, and/or the modification of the amino acid sequence of the molecule. In addition, a variety of different chemical modifications of the naturally occurring polypeptide can be used, either with or without modifications to the amino acid sequence or size of the molecule. Such chemical modifications can, for example, include the incorporation of modified or non-natural amino acids or non-amino acid moieties during synthesis of the peptide chain, or the post-synthesis modification of incorporated chain moieties.

The oligopeptides of this invention can be synthesized chemically or through an appropriate gene expression system. Synthetic peptides can include both naturally occurring amino acids and laboratory synthesized, modified amino acids.

Also provided herein are functional derivatives of anti-microbial proteins or polypeptides. By "functional derivative" is meant a "chemical derivative," "fragment," "variant," "chimera," or "hybrid" of the polypeptide or protein, which terms are defined below. A functional derivative retains at least a portion of the function of the protein, for example reactivity with a specific antibody, enzymatic activity or binding activity.

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A "chemical derivative" of the complex contains additional chemical moieties not normally a part of the protein or peptide. Such moieties may improve the molecule's solubility, absorption, biological half-life, and the like. The moieties may alternatively decrease the toxicity of the molecule, eliminate or attenuate any undesirable side effect of the molecule, and the like. Moieties capable of mediating such effects are disclosed in Alfonso and Gennaro (1995). Procedures for coupling such moieties to a molecule are well known in the art. Covalent modifications of the protein or peptides are included within the scope of this invention. Such modifications may be introduced into the molecule by reacting targeted amino acid residues of the peptide with an organic derivatizing agent that is capable of reacting with selected side chains or terminal residues, as described below.

Cysteinyl residues most commonly are reacted with alpha-haloacetates (and corresponding amines), such as chloroacetic acid or chloroacetamide, to give carboxymethyl or carboxyamidomethyl derivatives. Cysteinyl residues also are derivatized by reaction with bromotrifluoroacetone, chloroacetyl phosphate, N-alkylmaleimides, 3-nitro-2-pyridyl disulfide, methyl 2-pyridyl disulfide, p-chloromercuribenzoate, 2-chloromercuri-4-nitrophenol, or chloro-7-nitrobenzo-2-oxa-1,3-diazole.

Histidyl residues are derivatized by reaction with diethylprocarbonate at pH 5.5-7.0 because this agent is relatively specific for the histidyl side chain. Parabromophenacyl bromide also is useful; the reaction is preferably performed in 0.1 M sodium cacodylate at pH 6.0.

Lysinyl and amino terminal residues are reacted with succinic or other carboxylic acid anhydrides. Derivatization with these agents has the effect of reversing the charge of the lysinyl residues. Other suitable reagents for derivatizing primary amine- containing residues include imidoesters such as methyl picolinimidate; pyridoxal phosphate; pyridoxal; chloroborohydride;

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trinitrobenzenesulfonic acid; O-methylisourea; 2,4 pentanedione; and transaminase-catalyzed reaction with glyoxylate.

Arginyl residues are modified by reaction with one or several conventional reagents, among them phenylglyoxal, 2,3-butanedione, 1,2-cyclohexanedione, and ninhydrin. Derivatization of arginine residues requires that the reaction be performed in alkaline conditions because of the high  $pK_a$  of the guanidine functional group. Furthermore, these reagents may react with the groups of lysine as well as the arginine alpha-amino group.

Tyrosyl residues are well-known targets of modification for introduction of spectral labels by reaction with aromatic diazonium compounds or tetranitromethane. Most commonly, N-acetylimidizol and tetranitromethane are used to form O-acetyl tyrosyl species and 3-nitro derivatives, respectively.

Carboxyl side groups (aspartyl or glutamyl) are selectively modified by reaction carbodiimide (R'-N-C-N-R') such as 1-cyclohexyl-3-(2-morpholinyl(4-ethyl) carbodiimide or 1-ethyl-3-(4-azonia-4,4-dimethylpentyl) carbodiimide. Furthermore, aspartyl and glutamyl residues are converted to asparaginyl and glutaminyl residues by reaction with ammonium ions.

Glutaminyl and asparaginyl residues are frequently deamidated to the corresponding glutamyl and aspartyl residues. Alternatively, these residues are deamidated under mildly acidic conditions. Either form of these residues falls within the scope of this invention.

Derivatization with bifunctional agents is useful, for example, for cross-linking component peptides to each other or the complex to a water-insoluble support matrix or to other macromolecular carriers. Commonly used cross-linking agents include, for example, 1,1-bis (diazoacetyl)-2-phenylethane, glutaraldehyde, N-hydroxysuccinimide esters, for example, esters with 4-azidosalicylic acid, homobifunctional imidoesters, including disuccinimidyl esters such as 3,3'-dithiobis(succinimidylpropionate), and bifunctional maleimides such as bis-N-maleimido-1,8-octane. Derivatizing agents such as methyl-3-[p-azidophenyl) dithiolpropioimidate yield photoactivatable intermediates that are capable of forming crosslinks in the presence of light. Alternatively, reactive water-insoluble matrices such as cyanogen bromide-activated carbohydrates and the reactive substrates described in U.S. Patent Nos. 3,969,287; 3,691,016; 4,195,128; 4,247,642; 4,229,537; and 4,330,440 are employed for protein immobilization.

Other modifications include hydroxylation of proline and lysine, – phosphorylation of hydroxyl groups of seryl or threonyl residues, methylation of the alpha-amino groups of lysine, arginine, and histidine side chains (Creighton, T.E.,

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Proteins: Structure and Molecular Properties, W.H. Freeman & Co., San Francisco, pp. 79-86 (1983)), acetylation of the N-terminal amine, and, in some instances, amidation of the C-terminal carboxyl groups.

Such derivatized moieties may improve the stability, solubility, absorption, biological half life, and the like. The moieties may alternatively eliminate or attenuate any undesirable side effect of the protein complex. Moieties capable of mediating such effects are disclosed, for example, in Alfonso and Gennaro (1995).

The term "fragment" is used to indicate a polypeptide derived from the amino acid sequence of the protein or polypeptide having a length less than the full-length polypeptide from which it has been derived. Such a fragment may, for example, be produced by proteolytic cleavage of the full-length protein. Preferably, the fragment is obtained recombinantly by appropriately modifying the DNA sequence encoding the proteins to delete one or more amino acids at one or more sites of the C-terminus, N-terminus, and/or within the native sequence.

Another functional derivative intended to be within the scope of the present invention is a "variant" polypeptide that either lacks one or more amino acids or contains additional or substituted amino acids relative to the native polypeptide. The variant may be derived from a naturally occurring polypeptide by appropriately modifying the protein DNA coding sequence to add, remove, and/or to modify codons for one or more amino acids at one or more sites of the C-terminus, N-terminus, and/or within the native sequence.

A functional derivative of a protein or polypeptide with deleted, inserted and/or substituted amino acid residues may be prepared using standard techniques well-known to those of ordinary skill in the art. For example, the modified components of the functional derivatives may be produced using site-directed mutagenesis techniques (as exemplified by Adelman et al., 1983, *DNA* 2:183; Sambrook et al., 1989) wherein nucleotides in the DNA coding sequence are modified such that a modified coding sequence is produced, and thereafter expressing this recombinant DNA in a prokaryotic or eukaryotic host cell, using techniques such as those described above. Alternatively, components of functional derivatives of complexes with amino acid deletions, insertions and/or substitutions may be conveniently prepared by direct chemical synthesis, using methods well-known in the art.

Insofar as other anti-microbial inhibitor compounds identified by the invention described herein may not be peptidal in nature, other chemical techniques exist to allow their suitable modification, as well, and according the desirable principles discussed above.

## Administration and Pharmaceutical Compositions

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For the therapeutic and prophylactic treatment of infection, the preferred method of preparation or administration of anti-microbial compounds will generally vary depending on the precise identity and nature of the anti-microbial being delivered. Thus, those skilled in the art will understand that administration methods known in the art will also be appropriate for the compounds of this invention.

The particularly desired anti-microbial can be administered to a patient either by itself, or in pharmaceutical compositions where it is mixed with suitable carriers or excipient(s). In treating an infection, a therapeutically effective amount of an agent or agents is administered. A therapeutically effective dose refers to that amount of the compound that results in amelioration of one or more symptoms of bacterial infection and/or a prolongation of patient survival or patient comfort.

Toxicity, therapeutic and prophylactic efficacy of anti-microbials can be determined by standard pharmaceutical procedures in cell cultures and/or experimental organisms such as animals, e.g., for determining the LD<sub>50</sub> (the dose lethal to 50% of the population) and the ED<sub>50</sub> (the dose therapeutically effective in 50% of the population). The dose ratio between toxic and therapeutic effects is the therapeutic index and it can be expressed as the ratio LD<sub>50</sub>/ED<sub>50</sub>. Compounds that exhibit large therapeutic indices are preferred. The data obtained from these cell culture assays and animal studies can be used in formulating a range of dosage for use in humans. The dosage of such compounds lies preferably within a range of circulating concentrations that include the ED<sub>50</sub> with little or no toxicity. The dosage may vary within this range depending upon the dosage form employed and the route of administration utilized.

For any compound identified and used in the method of the invention, the therapeutically effective dose can be estimated initially from cell culture assays. Such information can be used to more accurately determine useful doses in organisms such as plants and animals, preferably mammals, and most preferably humans. Levels in plasma may be measured, for example, by HPLC or other means appropriate for detection of the particular compound.

The exact formulation, route of administration and dosage can be chosen by the individual physician in view of the patient's condition (see e.g. Fingl et. al., in The Pharmacological Basis of Therapeutics, 1975, Ch. 1 p.1).

It should be noted that the attending physician would know how and when to terminate, interrupt, or adjust administration due to toxicity, organ dysfunction, or other systemic malady. Conversely, the attending physician would also know to adjust treatment to higher levels if the clinical response were not adequate (precluding

69

toxicity). The magnitude of an administered dose in the management of the disorder of interest will vary with the severity of the condition to be treated and the route of administration. The severity of the condition may, for example, be evaluated, in part, by standard prognostic evaluation methods. Further, the dose and perhaps dose frequency, will also vary according to the age, body weight, and response of the individual patient. A program comparable to that discussed above also may be used in veterinary or phyto medicine.

Depending on the specific infection target being treated and the method selected, such agents may be formulated and administered systemically or locally, i.e., topically. Techniques for formulation and administration may be found in Alfonso and Gennaro (1995). Suitable routes may include, for example, oral, rectal, transdermal, vaginal, transmucosal, intestinal, parenteral, intramuscular, subcutaneous, or intramedullary injections, as well as intrathecal, intravenous, or intraperitoneal injections.

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For injection, the agents of the invention may be formulated in aqueous solutions, preferably in physiologically compatible buffers such as Hanks' solution, Ringer's solution, or physiological saline buffer. For transmucosal administration, penetrants appropriate to the barrier to be permeated are used in the formulation. Such penetrants are generally known in the art.

Use of pharmaceutically acceptable carriers to formulate identified antimicrobials of the present invention into dosages suitable for systemic administration is within the scope of the invention. With proper choice of carrier and suitable manufacturing practice, the compositions of the present invention, in particular those formulated as solutions, may be administered parenterally, such as by intravenous injection. Appropriate compounds can be formulated readily using pharmaceutically acceptable carriers well known in the art into dosages suitable for oral administration. Such carriers enable the compounds of the invention to be formulated as tablets, pills, capsules, liquids, gels, syrups, slurries, suspensions and the like, for oral ingestion by a patient to be treated.

Agents intended to be administered intracellularly may be administered using techniques well known to those of ordinary skill in the art. For example, such agents may be encapsulated into liposomes, then administered as described above.

Liposomes are spherical lipid bilayers with aqueous interiors. All molecules present in an aqueous solution at the time of liposome formation are incorporated into the aqueous interior. The liposomal contents are both protected from the external microenvironment and, because liposomes fuse with cell membranes, are efficiently

70

delivered into the cell cytoplasm. Additionally, due to their hydrophobicity, small organic molecules may be directly administered intracellularly.

Pharmaceutical compositions suitable for use in the present invention include compositions wherein the active ingredients are contained in an effective amount to achieve the intended purpose. Determination of the effective amounts is well within the capability of those skilled in the art.

In addition to the active ingredients, these pharmaceutical compositions may contain suitable pharmaceutically acceptable carriers comprising excipients and auxiliaries which facilitate processing of the active compounds into preparations which can be used pharmaceutically. The preparations formulated for oral administration may be in the form of tablets, dragees, capsules, or solutions, including those formulated for delayed release or only to be released when the pharmaceutical reaches the small or large intestine.

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The pharmaceutical compositions of the present invention may be manufactured in a manner that is itself known, e.g., by means of conventional mixing, dissolving, granulating, dragee-making, levitating, emulsifying, encapsulating, entrapping or lyophilizing processes.

Pharmaceutical formulations for parenteral administration include aqueous solutions of the active anti-microbial compounds in water-soluble form.

Alternatively, suspensions of the active compounds may be prepared as appropriate oily injection suspensions. Suitable lipophilic solvents or vehicles include fatty oils such as sesame oil, or synthetic fatty acid esters, such as ethyl oleate or triglycerides, or liposomes. Aqueous injection suspensions may contain substances which increase the viscosity of the suspension, such as sodium carboxymethyl cellulose, sorbitol, or dextran. Optionally, the suspension may also contain suitable stabilizers or agents which increase the solubility of the compounds to allow for the preparation of highly concentrated solutions.

Pharmaceutical preparations for oral use can be obtained by combining the active compounds with solid excipient, optionally grinding a resulting mixture, and processing the mixture of granules, after adding suitable auxiliaries, if desired, to obtain tablets or dragee cores. Suitable excipients are, in particular, fillers such as sugars, including lactose, sucrose, mannitol, or sorbitol; cellulose preparations such as, for example, maize starch, wheat starch, rice starch, potato starch, gelatin, gum tragacanth, methyl cellulose, hydroxypropylmethyl-cellulose, sodium carboxymethylcellulose, and/or polyvinylpyrrolidone (PVP). If desired, disintegrating agents may be added, such as the cross-linked polyvinyl pyrrolidone, agar, or alginic acid or a salt thereof such as sodium alginate.

WO 00/32825 PCT/IB99/02040

71

Dragee cores are provided with suitable coatings. For this purpose, concentrated sugar solutions may be used, which may optionally contain gum arabic, talc, polyvinyl pyrrolidone, carbopol gel, polyethylene glycol, and/or titanium dioxide, lacquer solutions, and suitable organic solvents or solvent mixtures.

Dyestuffs or pigments may be added to the tablets or dragee coatings for identification

Dyestuits or pigments may be added to the tablets or dragee coatings for identification or to characterize different combinations of active compound doses.

Pharmaceutical preparations which can be used orally include push-fit

Pharmaceutical preparations which can be used orally include push-fit capsules made of gelatin, as well as soft, sealed capsules made of gelatin and a plasticizer, such as glycerol or sorbitol. The push-fit capsules can contain the active ingredients in admixture with filler such as lactose, binders such as starches, and/or lubricants such as talc or magnesium stearate and, optionally, stabilizers. In soft capsules, the active compounds may be dissolved or suspended in suitable liquids, such as fatty oils, liquid paraffin, or liquid polyethylene glycols. In addition, stabilizers may be added.

The above methodologies may be employed either actively or prophylactically against an infection of interest.

## Computer-related Aspects and Embodiments

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In addition to the provision of compounds as chemical entities, nucleotide sequences, or fragments thereof at least 95%, preferably at least 97%, more preferably at least 99%, and most preferably at least 99.9% identical to phage inhibitor sequences can also be provided in a variety of additional media to facilitate various uses.

Thus, as used in this section, "provided" refers to an article of manufacture, rather than an actual nucleic acid molecule, which contains a nucleotide sequence of the present invention; e.g., a nucleotide sequence of an exemplary bacteriophage or a sequence encoding a bacterial target or a fragment thereof, preferably a nucleotide sequence at least 95%, more preferably at least 99% and most preferably at least 99.9% identical to such a bacteriophage or bacterial sequence, for example, to a polynucleotide of an unsequenced phage listed in Table 1, preferably of bacteriophage 77 (S. aureus host) or bacteriophage 3A (S. aureus host) or bacteriophage 96 (S. aureus host). Such an article provides a large portion of the particular bacteriophage genome or bacterial gene and parts thereof (e.g., a bacteriophage open reading frame (ORF)) in a form which allows a skilled artisan to examine and/or analyze the sequence using means not directly applicable to examining the actual genome or gene or subset thereof as it exists in nature or in purified form as a chemical entity.

In one application of this aspect, a nucleotide sequence of the present invention can be recorded on computer readable media. As used herein, "computer

WO 00/32825 PCT/IB99/02040

72

readable media" refers to any medium that can be read and accessed directly by a computer. Such media include, but are not limited to: magnetic storage media, such as floppy discs, hard disc storage medium, magnetic tape; optical storage media such as CD-ROM; electrical storage media such as RAM and ROM; and hybrids of these categories, such as magnetic/optical storage media. A skilled artisan can readily appreciate how any of the presently known computer readable mediums can be used to create an article of manufacture which includes one or more computer readable media having recorded thereon a nucleotide sequence or sequences of the present invention. Likewise, it will be clear to those of skill how additional computer readable media that may be developed also can be used to create analogous manufactures having recorded thereon a nucleotide sequence of the present invention.

As used herein, "recorded" refers to a process for storing information on computer readable medium. A skilled artisan can readily adopt any of the presently known methods for recording information on computer readable medium to generate manufactures comprising the nucleotide sequence information of the present invention.

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A variety of data storage structures are available to a skilled artisan for creating a computer readable medium having recorded thereon a nucleotide sequence of the present invention. The choice of the data storage structure will generally be based on the means chosen to access the stored information. In addition, a variety of data processor programs and formats can be used to store the nucleotide sequence information of the present invention on computer readable medium. The sequence information can, for example, be presented in a word processing test file, formatted in commercially available software such as WordPerfect and Microsoft Word, or represented in the form of an ASCII file, stored in a database application, such as DB2, Sybase, Oracle, or the like. A skilled artisan can readily adapt any number of data processor structuring formats (e.g., text file or database) in order to obtain computer readable medium having recorded thereon the nucleotide sequence information of the present invention.

Computer software is publicly available which allows a skilled artisan to access sequence information provided in a computer readable medium. Thus, by providing in computer readable form a nucleotide sequence of an unsequenced bacteriophage, such as an exemplary bacteriophage listed in Table 1 or of a sequence encoding a bacterial target or a fragment thereof, preferably a nucleotide sequence at least 95%, more preferably at least 99% and most preferably at least 99.9% identical to such a bacteriophage or bacterial sequence, for example, to a polynucleotide of bacteriophage 77 (S. aureus host) or bacteriophage 3A (S. aureus host) bacteriophage

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96 (S. aureus host), bacteriophage 44AHJD (S. aureus host), bacteriophage Dp-1 (Streptococcus pneumoniae host), or bacteriophage 182 (Enterococcus host) the present invention enables the skilled artisan to routinely access the provided sequence information for a wide variety of purposes.

Those skilled in the art understand that software can implement a variety of different search or analysis software which implement sequence search and analysis algorithms, e.g., the BLAST (Altschul et al., J. Mol. Biol. 215:403410 (1990) and BLAZE (Brutlag et al., Comp. Chem 17:203-207 (1993)) search algorithms. For example, such search algorithms can be implemented on a Sybase system and used to identify open reading frames (ORFs) within the bacteriophage genome which contain homology to ORFs or proteins from other viruses, e.g, other bacteriophage, and other organisms, e.g., the host bacterium. Among the ORFs discussed herein are protein encoding fragments of the bacteriophage genomes which encode bacteria-inhibiting proteins or fragments.

The present invention further provides systems, particularly computer-based systems, which contain the sequence information described. Such systems are designed to identify, among other things, useful fragments of the bacteriophage genomes.

As used herein, "a computer-based system" refers to the hardware, software, and data storage media used to analyze the nucleotide sequence information of the present invention. The minimum hardware of the computer-based systems of the present invention comprises a central processing unit (CPU), input device, output device, and data storage medium or media. A skilled artisan will readily recognize that any of the currently available general purpose computer-based system are suitable for use in the present invention, as well as a variety of different specialized or dedicated computer-based systems.

As stated above, the computer-based systems of the present invention comprise data storage media having stored therein a nucleotide sequence of the present invention and the necessary hardware and software for supporting and implementing a search and/or analysis program.

As used herein, "data storage media" refers to memory which can store nucleotide sequence information of the present invention, or a memory access means which can access manufactures having recorded thereon the nucleotide sequence information of the present invention.

As used herein, "search program" refers to one or more programs which are implemented on the computer-based system to compare a target sequence or target structural motif with the sequence information stored within the data storage means.

WO 00/32825 PCT/IB99/02040

74

Search means are used to identify fragments or regions of the present gnomic sequences which match a particular target sequence or target motif. A variety of known algorithms are disclosed publicly and a variety of commercially available software for conducting search means are and can be used in the computer-based systems of the present invention. Examples of such software includes, but is not limited to, MacPattern (EMBL), BLASTN and BLASTX (NCBIA). A skilled artisan can readily recognize that any one of the available algorithms or implementing software packages for conducting homology searches and/or sequence analyses can be adapted for use in the present computer-based systems.

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As used herein in connection with sequence searches and analyses, a "target sequence" can be any DNA or amino acid sequence of six or more nucleotides or two or more amino acids. A skilled artisan can readily recognize that the longer a target sequence is, the less likely a target sequence will be present as a random occurrence in the database. Also, the target sequence length is preferably selected to include sequence corresponding to a biologically relevant portion of an encoded product, for example a region which is expected to be conserved across a range of source organisms. Preferably the sequence length of a target polypeptide sequence is from 5-100 amino acids, more preferably 7-50 or 7-100 amino acids, and still more preferably 10-80 or 10-100 amino acids. Preferably the sequence length of a target polynucleotide sequence is from 15-300 nucleotide residues, more preferably from 21-240 or 21-300, and still more preferably 30-150 or 30-300 nucleotide residues. However, it is well recognized that searches for commercially important fragments, such as sequence fragments involved in gene expression and protein processing, may be of shorter length. Likewise, it may be desirable to search and/or analyze longer sequences.

As used herein, "a target structural motif," or "target motif," refers to any rationally selected sequence or combination of sequences in which the sequence(s) are chosen based on a three-dimensional configuration which is formed upon the folding of the target motif. There are a variety of target motifs known in the art. Protein target motifs include, but are not limited to, enzymatic active sites and signal sequences. Nucleic acid target motifs include, but are not limited to promoter sequences, hairpin structures and inducible expression elements (protein binding sequences).

A variety of structural formats for the input and output devices can be used to input and output the information in the computer-based systems of the present invention. A preferred format for an output device ranks fragments of the bacteriophage or bacterial sequences possessing varying degrees of homology to the

WO 00/32825 PCT/IB99/02040

75

target sequence or target motif. Such presentation provides a skilled artisan with a ranking of sequences which contain various amounts of the target sequence or target motif and identifies the degree of homology contained in the identified fragment.

A variety of comparing methods and/or devices and/or formats can be used to compare a target sequence or target motif with the sequence stored in data storage media to identify sequence fragments of the bacteriophage or bacterium in question. One skilled in the art can readily recognize that any one of the publicly available homology search programs can be used as the search program for the computer-based systems of the present invention. Of course, suitable proprietary systems that may be known to those of skill, or later developed, also may be employed in this regard.

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Figure 6 provides a block diagram of a computer system illustrative of embodiments of this aspect of present invention. The computer system 102 includes a processor 106 connected to a bus 104. Also connected to the bus 104 are a main memory 108 (preferably implemented as random access memory, RAM) and a variety of secondary storage devices 110, such as a hard drive 112 and a removable medium storage device 114. The removable medium storage device114 may represent, for example, a floppy disk drive, a CD-ROM drive, a magnetic tape drive, etc. A removable storage medium 116 (such as a floppy disk, a compact disk, a magnetic tape, etc.) containing control logic and/or data recorded therein may be inserted into the removable medium storage device 114. The computer system 102 includes appropriate software for reading the control logic and/or the data from the removable medium storage device 114, once it is inserted into the removable medium storage device 114.

A nucleotide sequence of the present invention may be stored in a well-known manner in the main memory 108, any of the secondary storage devices 110, and/or a removable storage medium 116. During execution, software for accessing and processing the sequence (such as search tools, comparing tools, etc.) reside in main memory 108, in accordance with the requirements and operating parameters of the operating system, the hardware system and the software program or programs.

The data storage medium in which the sequence is embodied and the central processor need not be part of a single stand-alone computer, but may be separated so long as data transfer can occur. For example, the processor or processors being utilized for a search or analysis can be part of one general purpose computer, and the data storage medium can be part of a second general purpose computer connected to a network, or the data storage medium can be part of a network server. As another example the data storage medium can be part of a computer system or network accessible over telephone lines or other remote connection method.

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#### **EXAMPLES**

Example 1. Growth of Staph A bacteriophage 77 and purification of genomic DNA.

The Staphylococcus aureus propagating strain (PS 77; ATCC #27699) was used as a host to propagate its respective phage 77 (ATCC # 27699-B1). Two rounds of plaque purification of phage 77 were performed on soft agar essentially as described in Sambrook et al (1989). Briefly, the PS 77 strain was grown overnight at 37°C in Nutrient broth [NB: 0.3% Bacto beef extract, 0.5% Bacto peptone (Difco Laboratories) and 0.5% NaCl (w/v)]. The culture was then diluted 20x in NB and incubated at 37°C until the OD<sub>540</sub>= .2 (early log phase) with constant agitation. In order to obtain single plaques, phage 77 was subjected to 10-fold serial dilutions using phage buffer (1 mM MgSO<sub>4</sub>, 5 mM MgCl<sub>2</sub>, 80 mM NaCl and 0.1% Gelatin (w/v)) and 10 µl of each dilution was used to infect 0.5 ml of the cell suspension in the presence of 400 µg/ml CaCl<sub>2</sub>. After incubation of 15 min at room temperature (RT), 2 ml of melted soft agar kept at 45°C (NB supplemented with 0.6% agar) was added to the mixture and poured onto the surface of 100 mm nutrient agar plates (0.3% Bacto Beef extract, 0.5% Bacto peptone, 0.5% NaCl and 1.5% Bacto agar (w/v)). After overnight incubation at 30°C, a single plaque was isolated, resuspended in 1 ml of phage buffer by end over end rotation for 2 hrs at 20°C, and the phage suspension was diluted and used for a second infection as described above. After overnight incubation at 30°C, a single plaque was isolated and used as a stock.

The propagation procedure for bacteriophage 77 was modified from the agar layer method of Swanstörm and Adams (1951). Briefly, the PS 77 strain was grown to stationary phase overnight at 37°C in Nutrient broth. The culture was then diluted twenty-fold in NB and incubated at 37°C until the OD<sub>540</sub>= .2. The suspension (15x10<sup>7</sup> Bacteria) was then mixed with 15x10<sup>5</sup> plaque forming units (pfu) to give a ratio of 100-bacteria/phage particle in the presence of 400 μg/ml of CaCl<sub>2</sub>. After incubation for 15 min at 20°C, 7.5 ml of melted soft agar (NB plus 0.6% agar) were added to the mixture and poured onto the surface of 150 mm nutrient agar plates and incubated 16 hrs at 30°C. To collect the phage plate lysate, 20 ml of NB were added to each plate and the soft agar layer was collected by scrapping off with a clean microscope slide followed by shaking of the agar suspension for 5 min to break up the agar. The mixture was then centrifuged for 10 min at 4,000 RPM (2,830xg) in a JA-10 rotor—(Beckman) and the supernatant fluid (lysate) was collected and subjected to a treatment with 10 μg/ml of DNase I and RNase A for 30 min at 37°C. To precipitate the phage particles, the phage suspension was adjusted to 10% (w/v) PEG 8000 and

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0.5 M of NaCl followed by incubation at 4°C for 16 hrs. The phage was recovered by centrifugation at 4,000 rpm (3,500xg) for 20 min at 4°C on a GS-6R table top centrifuge (Beckman). The pellet was resuspended with 2 ml of phage buffer (1 mM MgSO<sub>4</sub>, 5 mM MgCl<sub>2</sub>, 80 mM NaCl and 0.1% Gelatin). The phage suspension was extracted with 1 volume of chloroform and further purified by centrifugation on a cesium chloride step gradient as described in Sambrook et al. (1989), using a TLS 55 rotor centrifuged in an Optima TLX ultracentrifuge (Beckman) for 2 h at 28,000 mm (67,000xg) at 4°C. Banded phage was collected and ultracentrifuged again on an isopycnic cesium chloride gradient (1.45 g/ml) at 40,000 rpm (64,000xg) for 24 h at 4°C using a TLV rotor (Beckman). The phage was harvested and dialyzed for 4 h at room temperature against 4 L of dialysis buffer consisting of 10 mM NaCl, 50 mM Tris-HCl [pH 8] and 10 mM MgCl<sub>2</sub>. Phage DNA was prepared from the phage suspension by adding 20 mM EDTA, 50 mg/ml Proteinase K and 0.5% SDS and incubating for 1 h at 65°C, followed by successive extractions with 1 volume of phenol, 1 volume of phenol-chloroform and 1 volume of chloroform. The DNA was then dialyzed overnight at 4°C against 4 L of TE (10 mM Tris pH 8.0, 1mM EDTA).

### Example 2. DNA sequencing of Bacteriophage 77 genome

Four micrograms of phage 77 DNA was diluted in 200 μl of TE (10 mM Tris, [pH 8.0], 1 mM EDTA) in a 1.5 ml eppendorf tube and sonication was performed (550 Sonic Dismembrator<sup>TM</sup>, Fisher Scientific). Samples were sonicated under an amplitude of 3 μm with bursts of 5 s spaced by 15 s cooling in ice/water for 3 to 4 cycles. The sonicated DNA was then size fractionated by electrophoresis on 1% agarose gels utilizing TAE (1 x TAE is: 40 mM Tris-acetate, 1 mM EDTA [pH 8.0]) as the running buffer. Fractions ranging from 1 to 2 kbp were excised from the agarose gel and purified using a commercial DNA extraction system according to the instructions of the manufacturer (Qiagen), with a final elution of 50 μl of 1 mM Tris (pH 8.5).

The ends of the sonicated DNA fragments were repaired with a combination of T4 DNA polymerase and the Klenow fragment of E. coli DNA polymerase I, as follows. Reactions were performed in a reaction mixture (final volume, 100 µl) containing sonicated phage DNA, 10 mM Tris-HCl [pH 8.0], 50 mM NaCl, 10 mM MgCl<sub>2</sub>, 1 mM DTT, 50 µg/ml BSA, 100 µM of each dNTP and 15 units of T4 DNA polymerase (New England Biolabs) for 20 min at 12°C followed by addition of 12.5 units of Klenow large fragment (New England Biolabs) for 15 min at room—temperature. The reaction was stopped by two phenol/chloroform extractions and the

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DNA was precipitated with ethanol and the final DNA pellet was resuspended in 20  $\mu$ l of H<sub>2</sub>O.

Blunt-ended DNA fragments were cloned by ligation directly into the *Hinc* II site of pKSII+ vector (New England Biolabs) dephosphorylated by treatment with calf intestinal alkaline phosphatase (New England Biolabs)-treated pKS II+ vector (Stratagene). A typical ligation reaction contained 100 ng of vector DNA, 2 to 5 μl of repaired sonicated phage DNA (50-100 ng) in a final volume of 20 μl containing 800 units of T4 DNA ligase (New England Biolabs) and was incubated overnight at 16°C. Transformation and selection of bacterial clones containing recombinant plasmids was performed in *E. coli* DH10β according to standard procedures (Sambrook et al., 1989).

Recombinant clones were picked from agar plates into 96-well plates containing 100 μl LB and 100 μg/ml ampicillin and incubated at 37°C. The presence of phage DNA insert was confirmed by PCR amplification using T3 and T7 primers flanking the *Hinc* II cloning site of the pKS II+ vector. PCR amplification of foreign insert was performed in a 15 μl reaction volume containing 10 mM Tris (pH 8.3), 50 mM KCl, 1.5 mM MgCl<sub>2</sub>, 0.02% gelatin, 1 μM primer, 187.5 μM each dNTP, and 0.75 units *Taq* polymerase (BRL). The thermocycling parameters were as follows: 2 min initial denaturation at 94°C for 2 min, followed by 20 cycles of 30 sec denaturation at 94°C, 30 sec annealing at 57°C, and 2 min extension at 72°C, followed by a single extension step at 72°C for 10 min. Clones with insert sizes of 1 to 2 kbp were selected and plasmid DNA was prepared from the selected clones using QIAprep<sup>TM</sup> spin miniprep kit (Qiagen).

The nucleotide sequence of the extremities of each recombinant clone was determined using an ABI 377-36 automated sequencer with two types of chemistry: ABI prism Big Dye™ primer or ABI prism Big Dye™ terminator cycle sequencing ready reaction kit (Applied Biosystems). To ensure co-linearity of the sequence data and the genome, all regions of phage genome were sequenced at least once from both directions on two separate clones. In areas that this criteria was not initially met, a sequencing primer was selected and phage DNA was used directly as sequencing template employing ABI prism Big Dye™ terminator cycle sequencing ready reaction kit.

Example 3. Bioinformatic management of primary nucleotide sequence from Phage 77.

Phage 77 sequence contigs were assembled using Sequencher<sup>TM</sup> 3.1 software (GeneCodes). To close contig gaps, sequencing primers were selected near the edge of

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the contigs. Phage DNA was used directly as sequencing template employing ABI prism BIG DYE™ terminator cycle sequencing ready reaction kit. The complete sequence of bacteriophage 77 is shown in Table 2.

A software program was developed and used on the assembled sequence of bacteriophage 77 to identify all putative ORFs larger than 33 codons. Other ORF identification software can also be utilized, preferably programs which allow alternative start codons. The software scans the primary nucleotide sequence starting at nucleotide #1 for an appropriate start codon. Three possible selections can be made for defining the nature of the start codon; I) selection of ATG, II) selection of ATG or GTG, and III) selection of either ATG, GTG, TTG, CTG, ATT, ATC, and ATA. This latter initiation codon set corresponds to the one reported by the NCBI (http://www.ncbi.nlm.nih.gov/htbin-post/Taxonomy/wprintgc?mode=c) for the bacterial genetic code.

When an appropriate start codon is encountered, a counting mechanism is employed to count the number of codons (groups of three nucleotides) between this start codon and the next stop codon downstream of it. If a threshold value of 33 is reached, or exceeded, then the sequence encompassed by these two codons (start and stop codons) is defined as an ORF. This procedure is repeated, each time starting at the next nucleotide following the previous stop codon found, in order to identify all the other putative ORFs. The scan is performed on all three reading frames of both DNA strands of the phage sequence.

Sequence homology (BLAST) searches for each ORF are then carried out using an implementation of BLAST programs, although any of a variety of different sequence comparison and matching programs can be utilized as known to those

- skilled in the art. Downloaded public databases used for sequence analysis include:
- i) non-redundant GenBank (ftp://ncbi.nlm.nih.gov/blast/db/nr.Z),
- ii) Swissprot (ftp://ncbi.nlm.nih.gov/blast/db/swissprot.Z);
- iii) vector (ftp://ncbi.nlm.nih.gov/blast/db/vector.Z);
- iv) pdbaa databases (ftp://ncbi.nlm.nih.gov/blast/db/pdbaa.Z);
- v) S. aureus NCTC 8325 (ftp://ftp.genome.ou.edu/pub/staph/staph-1k.fa);
  - vi) streptococcus pyogenes (ftp://ftp.genome.ou.edu/pub/strep/strep-1k.fa);
  - vii) Streptococcus pneumoniae
  - (ftp://ftp.tigr.org/pub/data/s\_pneumoniae/gsp.contigs.112197.Z);
  - viii) Mycobacterium tuberculosis CSU#9
- 35 (ftp://ftp.tigr.org/pub/data/m tuberculosis/TB 091097.Z) and
  - ix) pseudomonas aeruginosa (http://www.genome.washington.edu/pseudo/data.html).

The results of the homology searches performed on the ORFs is shown in Table 5.

# Example 4. Subcloning of Bacteriophage 77 ORFs into a Staph A inducible expression system.

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The shuttle vector pT0021, in which the firefly luciferase (lucFF) expression is controlled by the ars (arsenite) promoter/operator (Tauriainen et al., 1997), was modified in the following fashion. Two oligonucleotides corresponding to a short antigenic peptide derived from the heamaglutinin protein of influenza virus (HA epitope tag) were synthesized (Field et al., 1988). The sense strand HA tag sequence (with BamHI, SalI and HindIII cloning sites) is:

5'-gateccggtegaccaagettTACCCATACGACGTCCCAGACTACGCCAGCTGA-3' (where upper case letters denote the nucletotide sequence of the HA tag); the antisense strand HA tag sequence (with a HindIII cloning site) is:

5'-agctTCAGCTGGCGTAGTCTGGGACGTCGTATGGGTAaagcttggtcgaccgg-3' (where upper case letters denote the sequence of the HA tag). The two HA tag oligonucleotides were annealed and ligated into pT0021 vector which had been digested with BamHI and HindIII. This manipulation resulted in replacement of the lucFF gene by the HA tag. This modified shuttle vector containing the arsenite
 inducible promoter, the arsR gene, and HA tag was named pTHA. A diagram outlining our modification of pT0021 to generate pTHA is shown in Fig. 1A.

Each ORF, encoded by Bacteriophage 77, larger than 33 amino acids and having a Shine-Dalgarno sequence upstream of the initiation codon was selected for functional analysis for bacterial inhibition. In total, 98 ORFs were selected and screened as detailed below. A list of these is presented in Table 3. Each individual ORF, from initiation codon to last codon (excluding the stop codon), was amplified from phage genomic DNA using the polymerase chain reaction (PCR). For PCR amplification of ORFs, each sense strand primer targets the initiation codon and is preceded by a BamHI restriction site ("cgggatcc") and each antisense oligonucleotide targets the pentultimate codon (the one before the stop codon) of the ORF and is preceded by a Sal I restriction site ("gcgtcgaccg"). The PCR product of each ORF was gel purified and digested with BamHI and SalI. The digested PCR product was then gel purified using the Qiagen kit as described, ligated into BamHI and SaII digested pTHA vector, and used to transform E. coli bacterial strain DH10β(as described above). As a result of this manipulation, the HA tag is set inframe with the ORF and is positioned at the carboxy terminus of each ORF (pTHA/ORF clones). Recombinant pTHA/ORF clones were picked and their insert sizes were confirmed by PCR analysis

using primers flanking the cloning site. The names and sequences of the primers that were used for the PCR amplification were: HAF:

5'TATTATCCAAAACTTGAACA<sup>3</sup>; HAR: 5'CGGTGGTATATCCAGTGATT<sup>3</sup>'. The sequence integrity of cloned ORFs was verified directly by DNA sequencing using primers HAF and HAR. In cases where verification of ORF sequence could not be achieved by one pass with the sequencing primers, additional internal primers were selected and used for sequencing.

Staphylococcus aureus strain RN4220 (Kreiswirth et al., 1983) was used as a recipient for the expression of recombinant plasmids. Electoporation was performed essentially as previously described (Schenk and Laddaga, 1992). Selection of recombinant clones was performed on Luria-Broth agar (LB-agar) plates containing 30 µg/ml of kanamycin.

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For each ORF introduced in the pTHA plasmid, 3 independent transformants were isolated and used to individually inoculate cultures in 5 ml of TSB containing 30µg/ml kanamycin, followed by growth to saturation (16 hrs at 30°C). An aliquot of this stationary phase culture was used to generate a frozen glycerol stock of the transformant ( stored at - 80°C). The remaining culture was used for plasmid DNA extraction. Bacterial cells were harvested by centrifugation at 3000 x g at 22°C for 5 min. The pellet was resuspended in 200 µl 25% sucrose containing 25U/ml of lysostaphin and incubated for 15 min at 37°C. Then, 400µl of alkaline SDS solution (3% SDS, 0.2N NaOH) were added, well mixed and incubated for 7 min at room temperature. After the alkaline SDS treatment, 300µl of ice-cold 3M sodium acetate pH 4.8 were added, and the mix is immediately spun at 13000g for 15 min at room temperature. The supernatant was transferred to a new 1.5 ml conical centrifuge tube and 650µl of isopropanol (stored at room temperature) were added. The mix was then centrifuged at 13,000 x g for 5 min. The supernatant fluid was discarded, the pellet washed with 70% ethanol, and resuspended in 320 µl sterile distilled water.

The presence of individual phage 77 ORF DNA inserts in the plasmid was verified by PCR amplification using 1.5 µl transformant miniprep DNA in a PCR with primers flanking the cloning site of ORF in pTHA vector (HAF and HAR). The composition of the PCR reaction and the cycling parameters are identical to those employed for library screening described above.

Example 5. Functional assay for bacterial inhibitory activity of bacteriophage 77

ORFs.

The anti-microbial activity of individual phage 77 ORFs was monitored by two growth inhibitory assays, one on solid agar medium, the other in liquid medium.

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In general, Staphylococcus bacteria transformed with expression plasmids containing individual ORFs were grown in normal TSA medium and stored in 19% glycerol. At pre-determined times, arsenite was added to the culture to induce transcription of the phage 77 ORFs cloned immediately downstream from an arsenite-inducible promoter in the pTHA expression plasmid.

The effect of ORF induction on bacterial growth characteristics was then monitored and quantitated. The growth inhibition assay on solid medium was performed by streaking pTHA/ORF containing *S. aureus* transformant onto LB-Kn and TSA-Kn plates containing increasing concentrations of sodium arsenite (0; 2.5; 5; and 7.5 μM). Arsenite is used to induce the expression of cloned DNA in pTHA vector. In parallel, 3 μl of 1/10 and 1/100 dilutions of the frozen cultures of the pTHA/ORF transformants were spotted as single drops onto LB-Kn and TSA-Kn plates containing increasing concentration of sodium arsenite (0; 2.5; 5; and 7.5 μM). The plates were then incubated 16 hrs at 37°C, and the effect of arsenite-induced ORF expression on bacterial growth was monitored and quantitated by comparing the extent to that seen in control plates. As positive controls for growth inhibition,the *holin/lysin* genes of the *Sthaphylococcus aureus* phage Twort (Loessner et al., 1998) was subcloned into the pTHA *ars* inducible vector and used.

For the growth inhibition assay in liquid medium, stationary phase cultures were prepared by inoculating 2.5ml TSB-Kn with frozen S. aureus RN4220 transformants containing phage 77 ORFs cloned in pTHA vector followed by incubation for 16 hrs at 37°C. These cultures were then diluted 1/100 in the same medium, and the bacteria were allowed to grow for 2 hrs at 37°C to reach early log phase. 150 µl of such culture were then mixed with 2.35 ml TSB-Kn medium with or without arsenite (the final concentration of arsenite in the medium was 0 or 5 µM arsenite). After 3.5 hrs incubation at 37°C with shaking at 250 rpm, 100 µl of bacterial culture was removed from each tube for OD<sub>565</sub> measurement. Serial ten-fold dilutions of the culture in buffered saline solution (0.85% NaCl) were then spotted onto TSB-Kn plates. The plates were incubated at 37°C 16 hrs and the number of surviving colonies counted the following day. The growth inhibitory property of individual ORFs was then quantitated by comparing CFU numbers under normal or arsenite-induction conditions. A schematic flow of the inhibition analysis is shown in Fig. 3 (also applicable to inhibition analysis for the other phage and bacteria pointed out herein). Inhibition results are shown in Figures 4A-C.

Example 6: Itentification of Cecropin Signature Motif in Staphylococcus aureus

Bacteriophage 3A ORF

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The genome for S. aureus bacteriophage 3A was determined and the sequence was analyzed essentially as described for bacteriophage 77 in the examples above. Upon blast analysis of the identified open reading frames of phage 3A, the presence of an amino acid sequence corresponding to a cecropin signature motif was observed. This motif (WDGHKTLEK) is located at position aa 481-489. Cecropins were originally identified in proteins from the cecropia moth and are recognized as potent antibacterial proteins that constitute an important part of the cell-free immunity of insects. Cecropins are small proteins (31-39 amino acid residues) that are active against both Gram-positive and Gram-negative bacteria by disrupting the bacterial 10 membranes. Although the mechanisms by which the cecropons cause cell death are not fully understood, it is generally thought to involve channel formation and membrane destabilization.

The identification of a motif corresponding to a known inhibitor suggests that the product of ORF002 is also an inhibitory compound. Such inhibitory activity can be confirmed as described herein or by other methods known in the art. Confirmation of the inhibitory activity would indicate that the ORF product could serve as the basis for construction of mimetic compounds and other inhibitors directed to the target of the ORF002 product.

Boman & Hultmark, 1987, Ann. Rev. Microbiol. 41:103-126. Boman, 1991, Cell 65:205-207. Boman et al., 1991, Eur. J. Bioichem. 201:23-31. Wang et al., J. Biol. Chem. 273:27438-27448.

### Example 7. Growth of Staphylococcus aureus bacteriophage 44AHJD:

Staphylococcus aureus propagating strain (PS 44A) (Felix d'Herelle Reference Centre #HER 1101) was used as a host to propagate its respective phage 44AHJD (Felix d'Herelle Reference Centre #HER 101). Two rounds of plaque purification of phage 44AHJD were performed on soft agar essentially as described in Sambrook et al. (1989). Briefly, the Staphylococcus aureus PS strain was grown overnight at 37°C in Nutrient Broth [NB: 3 g Bacto Beef Extract, 5 g Bactopeptone per liter, (Difco Laboratories # 0003-17-8), supplemented with 0.5% NaCl]. The culture was then diluted 20 fold in NB and incubated at 37°C until an OD<sub>540</sub> of 0.2. In order to obtain single plaques, phage 44AHJD was subjected to 10-fold serial dilutions using the phage buffer (1 mM MgSO<sub>4</sub>, 5 mM MgCl<sub>2</sub>, 80 mM NaCl and 0.1% Gelatin) and 10 µl were used to infect 0.5 ml of the cell suspension in the presence of 400  $\mu$ g/ml of

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CaCl<sub>2</sub>. After incubation of 15 min at room temperature, 2 ml of melted soft agar (NB supplemented with 0.6% of agar) were added to the mixture and poured onto the surface of 100 mm nutrient agar plates (3 g Bacto Beef extract, 5 g Bactopeptone, 0.5% NaCl and 15 g of Bacto agar per liter (Difco Laboratories # 0001-17-0). After overnight incubation at 37°C, a single plaque was isolated, resuspended in 1ml of phage buffer by end over end rotation for 2 h at room temperature and the phage suspension was diluted and used for a second infection as described above. After overnight incubation at 37°C, a single plaque was isolated and used as a stock.

Large scale purification of bacteriophage and preparation of phage DNA was as follows.

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The propagation method was carried out by using the agar layer method described by Swanstorm and Adams (1951). Briefly, the PS 44A strain was grown to stationary phase overnight at 37°C in Nutrient Broth. The culture was then diluted 20x in NB and incubated at 37°C until the  $A_{540}$ = 0.2. The suspension (15x10<sup>7</sup> Bacteria) was then mixed with 15x10<sup>5</sup> phage particles to give a ratio of 100-bacteria/phage particle in the presence of 400 µg/ml of CaCl<sub>2</sub>. After incubation of 15 min at room temperature, 7.5 ml of melted soft agar were added to the mixture and poured onto the surface of 150 mm nutrient agar plates and incubated overnight at 37°C. To collect the lysate, 20 ml of NB were added to each plate and the soft agar layer was collected by scrapping off with a clean microscope slide and shaken vigorously for 5 min to break up the agar. The mixture was then centrifuged for 10 min at 4,000 rpm (2,830 xg) using a JA-10 rotor (Beckman) and the supernatant (lysate) is collected and subjected to a treatment with 10 µg/ml of DNase I and RNase A for 30 min at 37°C. To precipitate the phage particles, 10% (w/v) of PEG 8000 and 0.5 M of NaCl were added to the lysate and the mixture was incubated on ice for 16 h. The phage was recovered by centrifugation at 4,000 rpm (3,500 xg) for 20 min at 4°C on a GS-6R table top centrifuge (Beckman).

The pellet was resuspended with 2 ml of phage buffer (1 mM MgSO<sub>4</sub>, 5 mM MgCl<sub>2</sub>, 80 mM NaCl and 0.1% Gelatin). The phage suspension was extracted with 1 volume of chloroform and further purified by centrifugation on a preformed cesium chloride step gradient as described in Sambrook *et al.* (1989), using a TLS 55 rotor and centrifuged in an Optima TLX ultracentrifuge (Beckman) for 2 h at 28,000 rpm (67,000 xg) at 4°C. Banded phage was collected and ultracentrifuged again on an

isopycnic cesium chloride gradient (1.45 g/ml) at 40,000 rpm (64,000 x g) for 24 h at 4°C using a TLV rotor (Beckman). The phage was harvested and dialyzed for 4 h at room temperature against 4 L of dialysis buffer consisting of 10 mM NaCl, 50 mM Tris-HCl [pH 8] and 10 mM MgCl<sub>2</sub>. Phage DNA was prepared from the phage suspension by adding 20 mM EDTA, 50 µg/ml Proteinase K and 0.5% SDS and incubating for 1 h at 65°C, followed by successive extractions with 1 volume of phenol, 1 volume of phenol-chloroform and 1 volume of chloroform. The DNA was then dialyzed overnight at 4°C against 4 L of TE (10 mM Tris-HCl [pH 8.0], 1mM EDTA).

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in 20 µl of H<sub>2</sub>O.

### Example 8. DNA sequencing of the Bacteriophage 44 AHJD genome.

Four mg of phage DNA was diluted in 200 µl of TE pH 8.0 in a 1.5 ml eppendorf tube and sonication was performed (550 Sonic Dismembrator, Fisher Scientific). Samples were sonicated under an amplitude of 3 µm with bursts of 5 s spaced by 15 s cooling in ice/water for 3 to 4 cycles and size fractionated on 1% agarose gels. The sonicated DNA was then size fractionated by gel electrophoresis. Fractions ranging from 1 to 2 kbp were excised from the agarose gel and purified using a coommercial DNA extraction system according to the instructions of the manufacturer (Qiagen) and eluted in 50 µl of 1mMTris-HCl [ pH 8.5].

The ends of the sonicated DNA fragments were repaired with a combination of T4 DNA polymearse and the Klenow fragment of *E. coli* DNA polymerase 1 as follows. Reactions were performed in a final volume of 100 µl containing DNA, 10 mM Tris-HCl pH 8.0, 50 mM NaCl, 10 mM MgCl<sub>2</sub>, 1 mM DTT, 5 µg BSA, 100 µM of each dNTP and 15 units of T4 DNA polymerase (New England Biolabs) for 20 min at 12°C followed by addition of 12.5 units of Klenow fragment (New England Biolabs) for 15 min at room temperature. The reaction was stopped by two phenol/chloroform extractions and the DNA was ethanol precipitated and resuspended

Cloning of the sonicated phage DNA into pKSII vector and transformation:

Blunt-ended DNA fragments were cloned by ligation directly into the *HincII* site of the pkSII vector (Stratagene) dephosphorylated with calf intestinal alkaline phosphatase (New England Biolabs). A typical reaction contained 100 ng of vector, 2

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kit.

to 5  $\mu$ l of repaired sonicated phage DNA (50-100 ng) in a final volume of 20  $\mu$ l containing 800 units of T4 DNA ligase (New England Biolabs) overnight at 16°C. Transformation and selection of positive clones was performed in the host strain DH10  $\beta$  of *E. coli* using ampicillin as a selective antibiotic as described in Sambrook et al. (1989).

Recombinant clones were picked from agar plates into 96-well plates containing 100 ml LB and 100 µg/ml ampicillin and incubated at 37°C. The presence

of phage DNA insert was confirmed by PCR amplification using T3 and T7 primers flanking the HincII cloning site of the pKS vector. PCR amplification of the potential foreign inserts was performed in a 15 µl reaction volume containing 10 mM Tris-HCl (pH 8.3), 50 mM KCl, 1.5 mM MgCl<sub>2</sub>, 0.02% gelatin, 1 mM primer, 187.5 μM each dNTP, and 0.75 units Tag polymerase (BRL). The thermocycling parameters were as follows: 2 min initial denaturation at 94°C for 2 min, followed by 20 cycles of 30 sec denaturation at 94°C, 30 sec annealing at 58C, and 2 min extension at 72°C, followed by a single extension step at 72°C for 10 min. Clones with insert sizes of 1 to 2 kbp were selected and plasmid DNA was prepared from the selected clones using the QIAprep™ spin miniprep kit (Qiagen). The nucleotide sequence of the extremities of each recombinant clone was determined using an ABI 377-36 automated sequencer with two types of chemistry: ABI prism BigDye<sup>™</sup> primer cycle sequencing (21M13 primer: #403055)(M13REV primer: #403056) or ABI prism BigDye™ terminator cycle sequencing ready reaction kit (Applied Biosystems; #4303152). To ensure co-linearity of the sequence data and the genome, all regions of the phage genome were sequenced at least once from both directions on two separate clones. In areas that this criteria was not initially met, a sequencing primer was selected and phage DNA was used directly as sequencing template employing ABI prism BigDye™ terminator cycle sequencing ready reaction

# Example 9. Bioinformatic management of primary nucleotide sequence.

Sequence contigs were assembled using Sequencher<sup>TM</sup> 3.1 software (GeneCodes). To close contig gaps, sequencing primers were selected near the edge of the contigs. Phage DNA was used directly as sequencing template employing ABI

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prism BigDye<sup>™</sup> terminator cycle sequencing ready reaction kit (Applied Biosystems; #4303152). The complete sequence of *Staphylococcus aureus* bacteriophage 44AHJD is shown in Table 16.

A software program was used on the assembled sequence of bacteriophage 44AHJD to identify all putative ORFs larger than 33 codons. The software scans the primary nucleotide sequence starting at nucleotide #1 for an appropriate start codon. Three possible selections can be made for defining the nature of the start codon; I) selection of ATG, II) selection of ATG or GTG, and III) selection of either ATG, GTG, TTG, CTG, ATT, ATC, and ATA. This latter initiation codon set corresponds to the one reported by the NCBI(http://www.ncbi.nlm.nih.gov/htbinpost/Taxonomy/wprintgc?mode=c) for the bacterial genetic code. When an appropriate start codon is encountered, a counting mechanism is employed to count the number of codons (groups of three nucleotides) between this start codon and the next stop codon downstream of it. If a threshold value of 33 is reached, or exceeded, then the sequence encompassed by these two codons is defined as an ORF. This procedure is repeated, each time starting at the next nucleotide following the previous stop codon found, in order to identify all the other putative ORFs. The scan is performed on all three reading frames of both DNA strands of the phage sequence. The predicted ORFs for bacteriophage 44AHJD are listed in Tables 17 & 18.

Sequence homology searches for each ORF were carried out using an implementation of blast programs. Downloaded public databases used for sequence analysis include:

- (i) non-redundant GenBank (ftp://ncbi.nlm.nih.gov/blast/db/nr.Z),
- ii) Swissprot (ftp://ncbi.nlm.nih.gov/blast/db/swissprot.Z);
- 25 iii) vector (ftp://ncbi.nlm.nih.gov/blast/db/vector.Z);
  - iv) pdbaa databases (ftp://ncbi.nlm.nih.gov/blast/db/pdbaa.Z);
  - V) Staphylococcus aureus NCTC 8325 (ftp://ftp.genome.ou.edu/pub/staph/staphlk.fa);
  - vi) Staphylococcus pyogenes (ftp://ftp.tigr.org/pub/data/s\_pneumoniae/gsp.contigs.1121
  - vii)PRODOM(ftp://ftp.toulouse.inra.fr/pub/prodom/current\_release/prodom99\_1.forbl ast.gz);
  - viii) DOMO (ftp://ftp.infobiogen.fr/pub/db/domo/);

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ix) TREMBL (ftp://www.expasy.ch/databases/sp tr nrdb/fasta/)

The results of the homology searches performed on the ORFs of bacteriophage 44AHJD are shown in Tables 19 & 20.

## 5 Example 10. Sub-Cloning of Bacteriophage 44 AHJD ORFs.

Expression preferably utilizes a shuttle expression vector which is arranged such that expression of the exogenous bacteriophage 44 AHJD ORF sequence is inducible. For example, the shuttle vector pT0021, in which the firefly luciferase (lucFF) expression is controlled by the ars (arsenite) promoter/operator (Tauriainen et al., 1997), can be modified in the following fashion. Two oligonucleotides corresponding to a short antigenic peptide derived from the heamaglutinin protein of influenza virus (HA epitope tag) were synthesized (Field et al., 1988). The sense strand HA tag sequence (with BamHI, SalI and HindIII cloning sites) is: 5'-gatcccggtcgaccaagcttTACCCATACGACGTCCCAGACTACGCCAGCTGA-3' (where upper case letters denote the nucletotide sequence of the HA tag); the antisense strand HA tag sequence (with a HindIII cloning site) is: 5'-agctTCAGCTGGCGTAGTCTGGGACGTCGTATGGGTAaagcttggtcgaccgg-3' (where upper case letters denote the sequence of the HA tag). The two HA tag oligonucleotides were annealed and ligated into pT0021 vector which had been digested with BamHI and HindIII. This manipulation resulted in replacement of the lucFF gene by the HA tag. This modified shuttle vector containing the arsenite inducible promoter, the arsR gene, and HA tag was named pTHA. A diagram outlining our modification of pT0021 to generate pTHA is shown in Fig. 1A (another userful vector construct is shown in Fig. 1B).

Each ORF, encoded by Bacteriophage 44 AHJD, larger than 33 amino acids and having a Shine-Dalgarno sequence upstream of the initiation codon can be selected for functional analysis for bacterial inhibition. Each individual ORF, from initiation codon to last codon (excluding the stop codon), can be amplified from phage genomic DNA using the polymerase chain reaction (PCR). For PCR amplification of ORFs, each sense strand primer targets the initiation codon and is preceded by a BamHI restriction site (<sup>5</sup>cgggatcc<sup>3</sup>) and each antisense oligonucleotide targets the pentultimate codon (the one before the stop codon) of the ORF and is preceded by a Sal I restriction site (<sup>5</sup>gcgtcgaccg<sup>3</sup>). The PCR product of each ORF can be gel

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purified and digested with BamHI and SalI. The digested PCR product can then be gel purified using the Qiagen kit as described, ligated into BamHI and SalI digested pTHA vector, and used to transform E. coli bacterial strain DH10β(as described above). As a result of this manipulation, the HA tag is set inframe with the ORF and is positioned at the carboxy terminus of each ORF (pTHA/ORF clones). Recombinant pTHA/ORF clones will be picked and their insert sizes were confirmed by PCR analysis using primers flanking the cloning site. The following primers can be used for PCR amplification: HAF: 5'TATTATCCAAAACTTGAACA'; HAR: 5'CGGTGGTATATCCAGTGATT'. The sequence integrity of cloned ORFs can be verified directly by DNA sequencing using primers HAF and HAR. In cases where verification of ORF sequence can not be achieved by one pass with the sequencing primers, additional internal primers will be selected and used for sequencing.

Staphylococcus aureus strain RN4220 (Kreiswirth et al., 1983) will be used as a recipient for the expression of recombinant plasmids. Electoporation will be performed essentially as previously described (Schenk and Laddaga, 1992). Selection of recombinant clones will be performed on Luria-Broth agar (LB-agar) plates containing 30 µg/ml of kanamycin.

Alternatively, a constitutive promoter can be used to drive expression of the introduced ORF, and compare cell growth to control bacterial cells containing the parental vector lacking any introduced phage ORF. Recombinant plasmids will be introduced into *Staphylococcus aureus* strain RN4220 (Kreiswirth et al., 1983) using electoporation as previously described (Schenk and Laddaga, 1992).

## Cloning of ORFs with a Shine-Dalgarno sequence

ORFs with a Shine-Dalgamo sequence are selected for functional analysis of bacterial killing. Each ORF, from initiation codon to last codon (excluding the stop codon), can be amplified by PCR from phage genomic DNA. For PCR amplification of ORFs, each sense strand primer starts at the initiation codon and is preceded by a restriction site and each antisense strand starts at the last codon (excluding the stop codon) and is preceded by a different restriction site. The PCR product of each ORF will be gel purified and digested with the restriction enzymes with sites contained on the PCR oligonucleotides. The digested PCR product is then gel purified using the Qiagen kit, ligated into the modified shuttle vector, and used to transform bacterial strain DH10. Recombinant clones are then picked and their insert sizes confirmed by

PCR analysis using primers flanking the cloning site as well as restriction digestion. The sequence fidelity of cloned ORFs can be verified by DNA sequencing using the same primers as used for PCR. In the cases that the verification of ORFs can not be achieved by one path of sequencing using primers flanking the cloning site internal primers can be selected and used for sequencing. Recombinant plasmids can be introduced into *Staphylococcus aureus* strain RN4220 (Kreiswirth et al., 1983) using electoporation as previously described (Schenk and Laddaga, 1992).

### Induction of gene expression from the ars promoter.

If an inducible promoter is used, e.g., the *ars* promoter, induction can be assessed, for example, in either of the two methods.

# 1. Screening on agar plates

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The functional identification of killer ORFs can be performed by spreading an aliquot of *S. aureus* transformed cells containing phage 44 AHJD ORFs onto agar plates containing different concentrations of sodium arsenite (0; 2.5; 5; and 7.5 µM). The plates are incubated overnight at 37°C, after which a growth inhibition of the ORF transformants on plates that contain arsenite are compared to plates without arsenite. 2. Quantification of growth inhibition in liquid medium

Cells containing different recombinant plasmids can be grown for overnight at 37°C in LB medium supplemented with the appropriate antibiotic selection. These are then diluted to the mid log phase (OD<sub>540</sub>=.2) with fresh media containing antibiotic and transferred to 96-well microtitration plates (100 μl/well). Inducer is then added at different final concentrations (ranging from 2.5 to 10 μM) and the culture incubated for an additional 2 hrs at 37°C. The effect of expression of the phage 44 AHJD ORFs on bacterial cell growth is then monitored by measuring the OD<sub>540</sub> and comparing the rate of growth to the culture not containing inducer. [As positive controls for growth inhibition, the *kilA* gene of phage lambda (Reisinger, GR., Rietsch, A., Lubitz, W. and Blasi, U. 1993 *Virology* #193: 1033-1036), and the *holin/lysin* genes of the *Sthaphylococcus aureus* phage Twort (Loessner, MJ., Gaeng, S., Wendlinger, G., Maier, SK. and Scherer, S. 1998. *FEMS Microbiology Letters* #162:265-274) can be subcloned into the *ars* inducible vector. An aliquot of the induced and uninduced culture can also be plated out on agar plates containing an appropriate antibiotic-selection but lacking inducer. Following incubation overnight at 37°C, the number of

colonies is counted. Any ORF showing bacteriostatic activity will show a lower, but detectable, number of colonies on the agar plates when grown in the presence of inducer as compared to when grown in the absence of inducer. Any ORF showing full bacteriocidal activity will show no colonies on the agar plates, when grown in the presence of inducer as compared to when grown in the absence of inducer.

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# Example 11. Growth of *Enterococcus* bacteriophage 182 and purification of genomic DNA.

The Enterococcus propagating strain (PS) (Enterococcus sp. Group D, Felix d'Herelle Reference Centre #HER 1080) was used as host to propagate its respective phage 182 (Felix d'Herelle Reference Centre #HER 80). Two rounds of plaque purification of phage 182 were performed on soft agar essentially as described in Sambrook et al. (1989). Briefly, the Enterococcus sp. PS strain was grown overnight at 37°C in Tryptic Soy Broth [TSB: 17 g Bacto tryptone, 3 g Bacto soytone, 2.5 g Bacto dextrose, 5 g Sodium chloride, and 2.5 g Dipotassium phosphate per liter (Difco Laboratories (#0370-17-3)]. The culture was then diluted 20 fold in TSB and incubated at 37°C until the OD<sub>540</sub>= 0.2 (early log phase) with constant agitation. In order to obtain single plaques, phage 182 was subjected to 10 fold serial dilutions using the phage buffer (1 mM MgSO<sub>4</sub>, 5 mM MgCl<sub>2</sub>, 80 mM NaCl and 0.1% Gelatin (w/v)) and 10 l of each dilution was used to infect 0.5 ml of the bacterial cell suspension. After incubation at 15 min at 37°C, 2 ml of melted soft agar (TSB supplemented with 0.6% agar) was added to the mixture and poured onto the surface of 100 mm Trytic Soy Agar plates [TSA: 15 g Tryptone peptone, 5 g Soytone peptone, 5 g Sodium chloride and 15 g of Agar per liter (Difco Laboratories #0369-17)]. After overnight incubation at 37°C, a single plaque was isolated, resuspended in 1 ml of phage buffer by end over end rotation for 2 hrs at room temperature, and the phage suspension was diluted and used for a second infection as described above. After overnight incubation at 37°C, a single plaque was isolated and used as a stock for all subsequent manipulations.

The propagation procedure for bacteriophage 182 was modified from the agar layer method of Swanstörm and Adams (1951). Briefly, the *Enterococcus* sp. PS strain was grown to stationary phase overnight at 37°C in TSB. The culture was then diluted 20 fold in TSB and incubated at 37°C until the  $A_{540}$ = 0.2. The suspension (15x10<sup>7</sup> Bacteria) was then mixed with 15x10<sup>5</sup> plaque forming units (pfu) to give a

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ratio of 100-bacteria/pfu. After incubation of 15 min at 37°C, 7.5 ml of melted soft agar (TSB plus 0.6% agar) were added to the mixture and poured onto the surface of 150 mm TSA plates and incubated 16 hrs at 37°C. To collect the plate lysate, 20 ml of TSB were added to each plate and the soft agar layer was collected by scrapping off with a clean microscope slide followed by vigorous shaking of the agar suspension for 5 min to break up the agar. The mixture was then centrifuged for 10 min at 4,000 rpm (2,830 xg) using a JA-10 rotor (Beckman) and the supernatant fluid (lysate) is collected and subjected to a treatment with 10 µg/ml of DNase I and RNase A for 30 min at 37°C. To precipitate the phage particles, the phage suspension was adjusted to 10% (w/v) of PEG 8000 and 0.5 M of NaCl followed by incubation at 4°C for 16 hrs. The phage was recovered by centrifugation at 4,000 rpm (3,500 xg) for 20 min at 4°C on a GS-6R table top centrifuge (Beckman). The pellet was resuspended with 2 ml of phage buffer (1 mM MgSO<sub>4</sub>, 5 mM MgCl<sub>2</sub>, 80 mM NaCl and 0.1% Gelatin). The phage suspension was extracted with 1 volume of chloroform and further purified by centrifugation on a cesium chloride step gradient as described in Sambrook et al. (1989), using a TLS 55 rotor and centrifuged in an Optima TLX ultracentrifuge (Beckman) for 2 hrs at 28,000 rpm (67,000 xg) at 4°C. Banded phage was collected and ultracentrifuged again on an isopycnic cesium chloride gradient (1.45 g/ml) at 40,000 rpm (64,000 xg) for 24 hrs at 4°C using a TLV rotor (Beckman). The phages were harvested and dialyzed for 4 hrs at room temperature against 4 L of dialysis buffer consisting of 10 mM NaCl, 50 mM Tris-HCl [pH 8] and 10 mM MgCl, Phage DNA was prepared from the phage suspension by adding 20 mM EDTA, 50 g/ml Proteinase K and 0.5% SDS and incubating for 1 hr at 65°C, followed by successive extractions with 1 volume of phenol, 1 volume of phenol-chloroform and 1 volume of chloroform. The DNA was then dialyzed overnight at 4°C against 4 L of TE (10 mM Tris-HCl [pH 8.0], 1mM EDTA).

## Example 12. DNA sequencing of the Bacteriophage 182 genome.

Four micrograms of phage DNA was diluted in 200 µl of TE (10 mM Tris, [pH 8.0], 1 mM EDTA) in a 1.5 ml eppendorf tube and sonication was performed (550 Sonic Dismembrator, Fisher Scientific). Samples were sonicated under an amplitude of 3 µm with bursts of 5 s spaced by 15 s cooling in ice/water for 3 to 4

cycles. The sonicated DNA was then size fractionated by electrophoresis on 1% agarose gels utilizing TAE (1 x TAE is: 40 mM Tris-acetate, 1 mM EDTA [pH 8.0]) as the running buffer. Fractions ranging from 1 to 2 kbp were excised from the agarose gel and purified using a commercial DNA extraction system according to the instructions of the manufacturer (Qiagen), with a final elution of 50 µl of 1 mM Tris [pH 8.5].

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The ends of the sonicated DNA fragments were repaired with a combination of T4 DNA polymerase and the Klenow fragment of E. coli DNA polymerase I, as follows. Reactions were performed in a reaction mixture (final volume, 100 μl) containing sonicated phage DNA, 10 mM Tris-HCl [pH 8.0], 50 mM NaCl, 10 mM MgCl<sub>2</sub>, 1 mM DTT, 50 μg/ml BSA, 100 μM of each dNTP and 15 units of T4 DNA polymerase (New England Biolabs) for 20 min at 12°C followed by addition of 12.5 units of the Klenow large fragment of DNA polymerase I(New England Biolabs) for 15 min at room temperature. The reaction was stopped by two phenol/chloroform extractions and the DNA was precipitated with ethanol and the final DNA pellet resuspended in 20 μl of H<sub>2</sub>O.

Blunt-ended DNA fragments were cloned by ligation directly into the *Hinc* II site of the pKSII+ vector (New England Biolabs) dephosphorylated by treatment with calf intestinal alkaline phosphatase (New England Biolabs). A typical ligation reaction contained 100 ng of vector DNA, 2 to 5 μl of repaired sonicated phage DNA (50-100 ng) in a final volume of 20 μl containing 800 units of T4 DNA ligase (New England Biolabs) and was incubated overnight at 16°C. Transformation and selection of bacterial clones containing recombinant plasmids was performed in *E. coli* DH10β according to standard procedures (Sambrook *et al.*, 1989).

Recombinant clones were picked from agar plates into 96-well plates containing 100 μl LB and 100 μg/ml ampicillin and incubated at 37°C. The presence of phage DNA insert was confirmed by PCR amplification using T3 and T7 primers flanking the Hinc II cloning site of the pKS vector. PCR amplification of the potential foreign inserts was performed in a 15 μl reaction volume containing 10 mM Tris (pH 8.3), 50 mM KCl, 1.5 mM MgCl<sub>2</sub>, 0.02% gelatin, 1 μM primer, 187.5 μM each dNTP, and 0.75 units *Taq* polymerase (BRL). The thermocycling parameters were as follows: 2 min initial denaturation at 94°C for 2 min, followed by 20 cycles of 30 sec

denaturation at 94°C, 30 sec annealing at 58°C, and 2 min extension at 72°C, followed by a single extension step at 72°C for 10 min. Clones with insert sizes of 1 to 2 kbp were selected and plasmid DNA was prepared from the selected clones using the QIAprep™ spin miniprep kit (Qiagen).

The nucleotide sequence of the extremities of each recombinant clone was determined using an ABI 377-36 automated sequencer with two types of chemistry: ABI prism Big Dye<sup>TM</sup> primer cycle sequencing (21M13 primer: #403055)(M13REV primer: #403056) or ABI prism Big Dye<sup>TM</sup> terminator cycle sequencing ready reaction kit (Applied Biosystems; #4303152). To ensure co-linearity of the sequence data and the genome, all regions of the phage genome were sequenced at least once from both directions on two separate clones. In areas that this criteria was not initially met, a sequencing primer was selected and phage DNA was used directly as sequencing template employing ABI prism BigDye<sup>TM</sup> terminator cycle sequencing ready reaction kit.

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### Example 13. Bioinformatic management of primary nucleotide sequence.

Sequence contigs were assembled using Sequencher<sup>TM</sup> 3.1 software (GeneCodes). To close contig gaps, sequencing primers were selected near the edge of the contigs. Phage DNA was used directly as sequencing template employing ABI prism BigDye<sup>TM</sup> terminator cycle sequencing ready reaction kit (Applied Biosystems; #4303152). The complete sequence of *Enterococcus* bacteriophage 182 is shown in Table 21.

A software program was used on the assembled sequence of bacteriophage 182 to identify all putative ORFs larger than 33 codons. The software scans the primary nucleotide sequence starting at nucleotide #1 for an appropriate start codon. Three possible selections can be made for defining the nature of the start codon; I) selection of ATG, II) selection of ATG or GTG, and III) selection of either ATG, GTG, TTG, CTG, ATT, ATC, and ATA. This latter initiation codon set corresponds to the one reported by the NCBI(http://www.ncbi.nlm.nih.gov/htbin-

post/Taxonomy/wprintgc?mode=c) for the bacterial genetic code. When an appropriate start codon is encountered, a counting mechanism is employed to count the number of codons (groups of three nucleotides) between this start codon and the

next stop codon downstream of it. If a threshold value of 33 is reached, or exceeded, then the sequence encompassed by these two codons is defined as an ORF. This procedure is repeated, each time starting at the next nucleotide following the previous stop codon found, in order to identify all the other putative ORFs. The scan is

- performed on all three reading frames of both DNA strands of the phage sequence. The predicted ORFs for bacteriophage 182 are listed in Tables 22 & 23. Sequence homology searches for each ORF were carried out using an implementation of BLAST programs. Downloaded public databases used for sequence analysis include:
- 10 (i) non-redundant GenBank (ftp://ncbi.nlm.nih.gov/blast/db/nr.Z),
  - ii) Swissprot (ftp://ncbi.nlm.nih.gov/blast/db/swissprot.Z);
  - iii) vector (ftp://ncbi.nlm.nih.gov/blast/db/vector.Z);
  - iv) pdbaa databases (ftp://ncbi.nlm.nih.gov/blast/db/pdbaa.Z);
  - v) staphylococcus aureus NCTC 8325 (ftp://ftp.genome.ou.edu/pub/staph/staph-
- 15 1k.fa);
  - vi) streptococcus pyrogenes

(ftp://ftp.tigr.org/pub/data/s\_pneumoniae/gsp.contigs.112197.Z);

vii) PRODOM

(ftp://ftp.toulouse.inra.fr/pub/prodom/current\_release/prodom99.1.forblast.gz);

- viii) DOMO (ftp://ftp.infobiogen.fr/pub/db/domo/);
  - ix) TREMBL (ftp://www.expasy.ch/databases/sp\_tr\_nrdb/fasta/)

The results of the homology searches performed on the ORFs of bacteriophage 182 are shown in Tables 24 & 26.

### 25 Example 14. Sub-Cloning of Bacteriophage 182 ORFs.

## Preparation of the shuttle expression vector

Expression preferably utilizes a shuttle expression vector which is arranged such that expression of the exogenous bacteriophage 182 ORF sequence is inducible. For example, the plasmid pND50 replicates in *E. coli*, *E. faecalis*, and *S. aureus* 

(Yamagishi, J., Kojima, T., Oyamada, Y., Fujimoto, K., Hattori, H., Nakamura, S., and Inoue, M. 1996. Antimocrob. Agents Chemother. 40, 1157-1163). This plasmid—can be modified by conventional techniques to insert the inducible arsenite promoter, derived from the shuttle vector pT0021, in which the firefly luciferase (lucFF)

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expression is controlled by the *ars* promoter/operator from a *S. aureus* plasmid (Tauriainen, S., Karp, M., Chang, W and Virta, M. (1997). Recombinant luminescent bacteria for measuring bioavailable arsenite and antimonite. *Appl. Environ. Microbiol.* 63:4456-4461). This modified shuttle vector will contain the *ars* promoter, *arsR* gene and a cloning site for introduction of individual phage ORFs downstream from a shine-delgarno sequence.

Other inducible regulatory sequences can be utilized instead of the arsenite-inducible system. An example is a nisin-inducible system The nisA promoter activity is dependent on the proteins NisR and NisK, which constitute a two-component signal transduction system that responds to the extracellular inducer nisin. The nisin sensitivity and inducer concentration required for maximal induction varies among the strains, but is functional in *Streptococcus pyogenes*, *Streptococcus agalactiae*, *Streptococcus pneumoniae*, *Enterococcus faecalis*, and *Bacillus subtilis*. Significant induction of the nisA promoter (10- to 60-fold induction) can be obtained in all of the species. A vector containing this promoter was published as Eichenbaum Z, Federle MJ, Marra D, de Vos WM, Kuipers OP, Kleerebezem M, and Scott JR (1998) *Appl Environ Microbiol* 64, 2763-2769. Other vectors, e.g., plasmids, can also be utilized which will allow replication and transciption in *Enterococcus*.

Alternatively, a constitutive promoter can be used (e.g., the β-lactamase promoter is constitutive in *E. faecalis* – see ref. 1) to drive expression of the introduced ORF, and compare cell growth to control bacterial cells containing the parental vector lacking any introduced phage ORF. Recombinant plasmids are introduced into *E. faecalis* strain FA2-2 by electroporation, as previously described (Yamagishi, J., Kojima, T., Oyamada, Y., Fujimoto, K., Hattori, H., Nakamura, S., and Inoue, M. 1996. *Antimicrob. Agents Chemother.* 40, 1157-1163).

#### Cloning of ORFs with a Shine-Dalgarno sequence

ORFs with a Shine-Dalgarno sequence are selected for functional analysis of bacterial killing. Each ORF, from initiation codon to last codon (excluding the stop codon), will be amplified by PCR from phage genomic DNA. For PCR amplification of ORFs, each sense strand primer starts at the initiation codon and is preceded by a restriction site and each antisense strand starts at the last codon (excluding the stop codon) and is preceded by a different restriction site. The PCR product of each ORF will be gel purified and digested with the restriction enzymes with sites contained on

the PCR oligonucleotides. The digested PCR product is then gel purified using the Qiagen kit, ligated into the modified shuttle vector, and used to transform bacterial strain DH10β. Recombinant clones are then picked and their insert sizes confirmed by PCR analysis using primers flanking the cloning site as well as restriction digestion.

The sequence fidelity of cloned ORFs will be verified by DNA sequencing using the same primers as used for PCR. In the cases that the verification of ORFs can not be achieved by one path of sequencing using primers flanking the cloning site internal primers will be selected and used for sequencing. Recombinant plasmids will be introduced into E. faecalis strain FA2-2 by electroporation, as previously described (Yamagishi, J., Kojima, T., Oyamada, Y., Fujimoto, K., Hattori, H., Nakamura, S., and Inoue, M. 1996. Antimicrob. Agents Chemother. 40, 1157-1163).

### Induction of gene expression from the ars promoter.

If an inducible promoter is used, e.g., the ars promoter, induction can be assessed, for example, in either of the two methods.

#### 15 1. Screening on agar plates

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The functional identification of killer ORFs can be performed by spreading an aliquot of E. faecalis transformed cells containing phage 182 ORF onto agar plates containing different concentrations of sodium arsenite (0; 2.5; 5; and 7.5  $\mu$ M). The plates are incubated overnight at 37°C, after which a growth inhibition of the ORF

transformants on plates that contain arsenite are compared to plates without arsenite.

2. Quantification of growth inhibition in liquid medium

Cells containing different recombinant plasmids can be grown for overnight at 37°C in LB medium supplemented with the appropriate antibiotic selection. These are then diluted to the mid log phase (OD<sub>540</sub>=.2) with fresh media containing antibiotic and transferred to 96-well microtitration plates (100 μl/well). Inducer is then added at different final concentrations (ranging from 2.5 to 10 μM) and the culture incubated for an additional 2 h at 37°C. The effect of expression of the phage 182 ORFs on bacterial cell growth is then monitored by measuring the OD<sub>540</sub> and comparing the rate of growth to the culture not containing inducer. As positive controls for growth inhibition, the *kilA* gene of phage lambda (Reisinger, GR., Rietsch, A., Lubitz, W. and Blasi, U. 1993 *Virology* #193: 1033-1036), and the *holin/lysin* genes of the *Sthaphylococcus aureus* phage Twort (Loessner, MJ., Gaeng, S., Wendlinger, G.,

Maier, SK. and Scherer, S. 1998. FEMS Microbiology Letters #162:265-274) were subcloned into the ars inducible vector. An aliquot of the induced and uninduced culture can also be plated out on agar plates containing an appropriate antibiotic selection but lacking inducer. Following incubation overnight at 37°C, the number of colonies is counted. Any ORF showing bacteriostatic activity will show a lower, but detectable, number of colonies on the agar plates when grown in the presence of inducer as compared to when grown in the absence of inducer. Any ORF showing bacteriocidal activity will show no colonies on the agar plates, when grown in the presence of inducer as compared to when grown in the absence of inducer.

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# Example 15. Growth of Streptococcus bacteriophage Dp-1 and purification of genomic DNA.

The Streptococcus pneumoniae R6 propagating strain (PS) (Tomasz, 1966) was used as host to propagate its respective phage Dp-1 (McDonnell et al., 1975). (Alternatively, Streptococcus (Diplococcus) pneumoniae R36A could be used. Strain R36A is available from ATCC as #11733 or 27336. Streptococcus pneumoniae is also available from Felix d'Herelle Reference Center in Quebec, Canada as catalog number HER 1054. Other S. pneumoniae strains are also available from ATCC.)

Two rounds of plaque purification of phage Dp-1 were performed on soft agar essentially as described in Sambrook et al. (1989). Briefly, the Streptococcus R6 PS strain was grown overnight at 37°C in K-Cat media [K-Cat: 10 g Bacto casitone, 5 g Bacto tryptone, 1 g Yeast extract, 5g Potassium chloride, 0.2% Glucose, 30mM Potassium phosphate buffer [pH 8] and 250,000 Units Catalase per liter (Boehringer Mannheim #10683600). The culture was then diluted 20 fold in K-CAT and

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incubated at 37°C until the OD<sub>540</sub>= 0.2 (early log phase) with constant agitation. In order to obtain single plaques, Dp-1 phage was subjected to 10-fold serial dilutions using the phage buffer (100 mM Tris-HCl [pH 7.5], 100 mM NaCl and 10 mM MgCl<sub>2</sub>)and 10 µl of each dilution was used to infect 0.5 ml of the cell suspension. After incubation of 15 min at 37°C, 2 ml of melted soft agar (K-CAT supplemented with 0.8% of agar) were added to the mixture and poured onto the surface of 100 mm K-CAT agar plates [K-CAT supplemented with 1.2 % of agar]. After solidification of the soft agar layer, an additional 5 ml of melted soft agar was added to visualize distinct plaques (Ronda et al., 1978). After overnight incubation at 37°C, a single plaque was isolated, resuspended in 1 ml of phage buffer by end over end rotation for 2 hrs at room temperature, and the phage suspension was diluted and used for a second infection as described above. After overnight incubation at 37°C, a single plaque was isolated and used as a stock for all subsequent manipulations.

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The propagation procedure for bacteriophage Dp-1 was modified from the agar layer method of Swanstörm and Adams (1951). Briefly, the R6 strain of Streptococcus pneumoniae was grown to stationary phase overnight at 37°C in K-CAT. The culture was then diluted 20 fold in K-CAT and incubated at 37°C until the  $OD_{540} = 0.2$ . The suspension (15x10<sup>7</sup> Bacteria) was then mixed with 15x10<sup>5</sup> plaque forming units (pfu) to give a ratio of 100-bacteria/pfu. After incubation of 15 min at 37°C, 7.5 ml of melted soft agar (K-CAT plus 0.8% agar) were added to the mixture and poured onto the surface of 150 mm K-CAT agar plates and incubated 16 hrs at 37°C. After solidification of the soft agar layer, 7.5 ml of melted soft agar were added to each plate. To collect the plate lysate, 20 ml of K-CAT media were added to each plate and the soft agar layers were collected by scrapping off with a clean microscope slide followed by vigorous shaking of the agar suspension for 5 min to break up the agar. The mixture was then centrifuged for 10 min at 4,000 rpm (2,830 xg) using a JA-10 rotor (Beckman) and the supernatant (lysate) was collected and subjected to a treatment with 10 µg/ml of DNase I and RNase A for 30 min at 37°C. To precipitate the phage particles, the phage suspension was adjusted to 10% (w/v) of PEG 8000 and 0.5 M of NaCl followed by incubation at 4°C for 16 hrs. The phage was recovered by centrifugation at 4,000 rpm (3,500 xg) for 20 min at 4°C on a GS-6R table top centrifuge (Beckman). The pellet was resuspended with 2 ml of phage buffer (100 mM Tris-HCl [pH 7.5], 100 mM NaCl and 10 mM MgCl<sub>2</sub>). The phage suspension was extracted with 1 volume of chloroform and further purified by centrifugation on a cesium chloride step gradient as described in Sambrook et al. (1989), using a TLS-55 rotor and centrifuged in an Optima TLX ultracentrifuge (Beckman) for 2 hrs at 28,000 rpm (67,000 xg) at 4°C. Banded phage was collected and ultracentrifuged again on an

isopycnic cesium chloride gradient (1.45 g/ml) at 40,000 rpm (64,000 xg) for 24 hrs at 4°C using a TLV rotor (Beckman). The phage was harvested and dialyzed for 4 hrs at room temperature against 4 L of dialysis buffer consisting of 10 mM NaCl, 50 mM Tris-HCl [pH 8] and 10 mM MgCl<sub>2</sub>. Phage DNA was prepared from the phage suspension by adding 20 mM EDTA, 50 µg/ml Proteinase K and 0.5% SDS and incubating for 1 hr at 65°C, followed by successive extractions with 1 volume of phenol, 1 volume of phenol-chloroform and 1 volume of chloroform. The DNA was then dialyzed overnight at 4°C against 4 L of TE (10 mM Tris-HCl [pH 8.0], 1mM EDTA).

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### Example 16. DNA sequencing of the Bacteriophage Dp-1 genome.

Four micrograms of phage DNA was diluted in 200 µl of TE (10 mM Tris, [pH 8.0], 1 mM EDTA) in a 1.5 ml eppendorf tube and sonication was performed (550 Sonic Dismembrator, Fisher Scientific). Samples were sonicated under an amplitude of 3 µm with bursts of 5 sec spaced by 15 sec cooling in ice/water for 3 to 4 cycles. The sonicated DNA was then size fractionated by electrophoresis on 1% agarose gels utilizing TAE (1 x TAE is: 40 mM Tris-acetate, 1 mM EDTA [pH 8.0]) as the running buffer. Fractions ranging from 1 to 2 kbp were excised from the agarose gel and purified using a commercial DNA extraction system according to the instructions of the manufacturer (Qiagen), with a final elution of 50 µl of 1 mM Tris [pH 8.5].

The ends of the sonicated DNA fragments were repaired with a combination of T4 DNA polymerase and the Klenow fragment of E. coli DNA polymerase I, as follows. Reactions were performed in a reaction mixture (final volume,  $100 \,\mu$ l) containing sonicated phage DNA,  $10 \, \text{mM}$  Tris-HCl [pH 8.0],  $50 \, \text{mM}$  NaCl,  $10 \, \text{mM}$  MgCl<sub>2</sub>,  $1 \, \text{mM}$  DTT,  $50 \, \mu \text{g/ml}$  BSA,  $100 \, \mu \text{M}$  of each dNTP and  $15 \, \text{units}$  of T4 DNA polymerase (New England Biolabs) for  $20 \, \text{min}$  at  $12 \, ^{\circ}\text{C}$  followed by addition of  $12.5 \, \text{units}$  of the Klenow large fragment of DNA polymerase I (New England Biolabs) for  $15 \, \text{min}$  at room temperature. The reaction was stopped by two phenol/chloroform extractions and the DNA was precipitated with ethanol and the final DNA pellet resuspended in  $20 \, \mu \text{l}$  of  $H_2O$ .

Blunt-ended DNA fragments were cloned by ligation directly into the *Hinc* II site of the pKSII+ vector (New England Biolabs) dephosphorylated by treatment with calf intestinal alkaline phosphatase (New England Biolabs). A typical ligation reaction contained 100 ng of vector DNA, 2 to 5 µl of repaired sonicated phage DNA (50-100 ng) in a final volume of 20 µl containing 800 units of T4 DNA ligase (New England Biolabs) and was incubated overnight at 16°C. Transformation and selection

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of bacterial clones containing recombinant plasmids was performed in *E. coli* DH10β according to standard procedures (Sambrook *et al.*, 1989).

Recombinant clones were picked from agar plates into 96-well plates containing 100 μl LB and 100 μg/ml ampicillin and incubated at 37°C. The presence of phage DNA insert was confirmed by PCR amplification using T3 and T7 primers flanking the *Hinc* II cloning site of the pKS vector. PCR amplification of the potential foreign inserts was performed in a 15 μl reaction volume containing 10 mM Tris (pH 8.3), 50 mM KCl, 1.5 mM MgCl<sub>2</sub>, 0.02% gelatin, 1 μM primer, 187.5 μM each dNTP, and 0.75 units *Taq* polymerase (BRL). The thermocycling parameters were as follows: 2 min initial denaturation at 94°C for 2 min, followed by 20 cycles of 30 sec denaturation at 94°C, 30 sec annealing at 58°C, and 2 min extension at 72°C, followed by a single extension step at 72°C for 10 min. Clones with insert sizes of 1 to 2 kbp were selected and plasmid DNA was prepared from the selected clones using the QIAprep<sup>TM</sup> spin miniprep kit (Qiagen).

The nucleotide sequence of the extremities of each recombinant clone was determined using an ABI 377-36 automated sequencer with two types of chemistry: ABI prism Big Dye<sup>TM</sup> primer cycle sequencing (21M13 primer: #403055)(M13REV primer: #403056) or ABI prism Big Dye<sup>TM</sup> terminator cycle sequencing ready reaction kit (Applied Biosystems; #4303152). To ensure co-linearity of the sequence data and the genome, all regions of the phage genome were sequenced at least once from both directions on two separate clones. In areas that this criteria was not initially met, a sequencing primer was selected and phage DNA was used directly as sequencing template employing ABI prism Big Dye<sup>TM</sup> terminator cycle sequencing ready reaction kit.

Example 17. Bioinformatic management of primary nucleotide sequence.

Sequence contigs were assembled using Sequencher<sup>™</sup> 3.1 software (GeneCodes). To close contig gaps, sequencing primers were selected near the edge of the contigs. Phage DNA was used directly as sequencing template employing ABI prism BigDye<sup>™</sup> terminator cycle sequencing ready reaction kit (Applied Biosystems; #4303152). The complete sequence of *Streptococcus* bacteriophage Dp-1 is shown in Table 28.

A software program was used on the assembled sequence of bacteriophage

Dp-1 to identify all putative ORFs larger than 33 codons. The software scans the
primary nucleotide sequence starting at nucleotide #1 for an appropriate start codon.

Three possible selections can be made for defining the nature of the start codon; I)
selection of ATG, II) selection of ATG or GTG, and III) selection of either ATG,

GTG, TTG, CTG, ATT, ATC, and ATA. This latter initiation codon set corresponds to the one reported by the NCBI(http://www.ncbi.nlm.nih.gov/htbin-post/Taxonomy/wprintgc?mode=c) for the bacterial genetic code. When an appropriate start codon is encountered, a counting mechanism is employed to count the number of codons (groups of three nucleotides) between this start codon and the next stop codon downstream of it. If a threshold value of 33 is reached, or exceeded, then the sequence encompassed by these two codons is defined as an ORF. This procedure is repeated, each time starting at the next nucleotide following the previous stop codon found, in order to identify all the other putative ORFs. The scan is performed on all three reading frames of both DNA strands of the phage sequence. The predicted ORFs for bacteriophage Dp-1 are listed in Tables 29 and 30, and Fig. 6.

Sequence homology searches for each ORF were carried out using an implementation of BLAST programs. Downloaded public databases used for sequence analysis include:

- (i) non-redundant GenBank (ftp://ncbi.nlm.nih.gov/blast/db/nr.Z),
- ii) Swissprot (ftp://ncbi.nlm.nih.gov/blast/db/swissprot.Z);
- iii) vector (ftp://ncbi.nlm.nih.gov/blast/db/vector.Z);
- iv) pdbaa databases (ftp://ncbi.nlm.nih.gov/blast/db/pdbaa.Z);
- v) staphylococcus aureus NCTC 8325
- 20 (ftp://ftp.genome.ou.edu/pub/staph/staph-1k.fa);
  - vi) streptococcus pyogenes

(ftp://ftp.tigr.org/pub/data/s pneumoniae/gsp.contigs.112197.Z);

vii) PRODOM

(ftp://ftp.toulouse.inra.fr/pub/prodom/current\_release/prodom99.1.forblast.gz);

viii) DOMO (ftp://ftp.infobiogen.fr/pub/db/domo/);

ix) TREMBL (ftp://www.expasy.ch/databases/sp tr nrdb/fasta/)

The results of the homology searches performed on the ORFs of bacteriophage Dp-1 are shown in Table 31.

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### Example 18. Sub-Cloning of Bacteriophage Dp-1 ORFs.

#### Preparation of the shuttle expression vector

Expression preferably utilizes a shuttle expression vector which is arranged such that expression of the exogenous bacteriophage Dp-1 ORF sequence is inducible.

For example, the plasmid pLSE4 replicates in *E. coli*, and *S. pneumoniae* (Diaz and Garcia, 1990). This plasmid can be modified by conventional techniques to insert the inducible arsenite promoter, derived from the shuttle vector pT0021, in which the

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firefly luciferase (*lucFF*) expression is controlled by the *ars* promoter/operator from a *S. aureus* plasmid (Tauriainen, S., Karp, M., Chang, W and Virta, M. (1997). Recombinant luminescent bacteria for measuring bioavailable arsenite and antimonite. *Appl. Environ. Microbiol.* 63:4456-4461). This modified shuttle vector will contain the *ars* promoter, *arsR* gene and a cloning site for introduction of individual phage ORFs downstream from a shine-dalgarno sequence.

Other inducible regulatory sequences can be utilized instead of the arsenite-inducible system. An example is a nisin-inducible system The nisA promoter activity is dependent on the proteins NisR and NisK, which constitute a two-component signal transduction system that responds to the extracellular inducer nisin. The nisin sensitivity and inducer concentration required for maximal induction varies among the strains, but is functional in *Streptococcus pyogenes*, *Streptococcus agalactiae*, *Streptococcus pneumoniae*, *Enterococcus faecalis*, and *Bacillus subtilis*. Significant induction of the nisA promoter (10- to 60-fold induction) can be obtained in all of the species. A vector containing this promoter was published as Eichenbaum Z, Federle MJ, Marra D, de Vos WM, Kuipers OP, Kleerebezem M, and Scott JR (1998) *Appl Environ Microbiol* 64, 2763-2769. Other vectors, e.g., plasmids, can also be utilized which will allow replication and transcription in *Streptococcus*.

Alternatively, a constitutive promoter can be used to drive expression of the introduced ORF, and compare cell growth to control bacterial cells containing the parental vector lacking any introduced phage ORF. Recombinant plasmids are introduced into *S. pneumoniae* R6 as previously described (Diaz and Garcia, 1990)

#### Cloning of ORFs with a Shine-Dalgamo sequence

ORFs with a Shine-Dalgarno sequence are selected for functional analysis of bacterial killing. Each ORF, from initiation codon to last codon (excluding the stop codon), will be amplified by PCR from phage genomic DNA. For PCR amplification of ORFs, each sense strand primer starts at the initiation codon and is preceded by a restriction site and each antisense strand starts at the last codon (excluding the stop codon) and is preceded by a different restriction site. The PCR product of each ORF will be gel purified and digested with the restriction enzymes with sites contained on the PCR oligonucleotides. The digested PCR product is then gel purified using the Qiagen kit, ligated into the modified shuttle vector, and used to transform bacterial strain DH10β. Recombinant clones are then picked and their insert sizes confirmed by PCR analysis using primers flanking the cloning site as well as restriction———digestion. The sequence fidelity of cloned ORFs will be verified by DNA sequencing using the same primers as used for PCR. In the cases that the verification of ORFs can not be achieved by one path of sequencing using primers flanking the cloning site

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internal primers will be selected and used for sequencing. Recombinant plasmids will be introduced into S. pneumoniae R6 as previously described (Diaz and Garcia, 1990). Induction of gene expression from the ars promoter.

If an inducible promoter is used, e.g., the ars promoter, induction can be assessed, for example, in either of the two methods.

### 1. Screening on agar plates

The functional identification of killer ORFs can be performed by spreading an aliquot of S. pneumoniae transformed cells containing phage Dp-1 ORFs onto agar plates containing different concentrations of sodium arsenite (0; 2.5; 5; and 7.5  $\mu$ M). The plates are incubated overnight at 37°C, after which a growth inhibition of the ORF transformants on plates that contain arsenite are compared to plates without arsenite.

## 2. Quantification of growth inhibition in liquid medium

Cells containing different recombinant plasmids can be grown for overnight at 37°C in LB medium supplemented with the appropriate antibiotic selection. These are then diluted to the mid log phase (OD<sub>540</sub>=.2) with fresh media containing antibiotic and transferred to 96-well microtitration plates (100 µl/well). Inducer is then added at different final concentrations (ranging from 2.5 to 10 µM) and the culture incubated for an additional 2 hrs at 37°C. The effect of expression of the phage Dp-1 ORFs on bacterial cell growth is then monitored by measuring the OD<sub>540</sub> and comparing the rate of growth to the culture not containing inducer. [As positive controls for growth inhibition, the kilA gene of phage lambda (Reisinger, GR., Rietsch, A., Lubitz, W. and Blasi, U. 1993 Virology #193: 1033-1036), and the holin/lysin genes of the Sthaphylococcus aureus phage Twort (Loessner, MJ., Gaeng, S., Wendlinger, G., Maier, SK. and Scherer, S. 1998. FEMS Microbiology Letters #162:265-274) can be 25 subcloned into the ars inducible vector. An aliquot of the induced and uninduced culture can also be plated out on agar plates containing an appropriate antibiotic selection but lacking inducer. Following incubation overnight at 37°C, the number of colonies is counted. Any ORF showing bacteriostatic activity will show a lower, but detectable, number of colonies on the agar plates when grown in the presence of inducer as compared to when grown in the absence of inducer. Any ORF showing full bacteriocidal activity will show no colonies on the agar plates, when grown in the presence of inducer as compared to when grown in the absence of inducer.

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All patents and publications mentioned in the specification are indicative of the levels of skill of those skilled in the art to which the invention pertains. All references cited in this disclosure are incorporated by reference to the same extent as if each reference had been incorporated by reference in its entirety individually.

One skilled in the art would readily appreciate that the present invention is well adapted to carry out the objects and obtain the ends and advantages mentioned, as well as those inherent therein. The specific methods and compositions described herein as presently representative of preferred embodiments are exemplary and are not intended as limitations on the scope of the invention. Changes therein and other uses will occur to those skilled in the art which are encompassed within the spirit of the invention are defined by the scope of the claims.

It will be readily apparent to one skilled in the art that varying substitutions and modifications may be made to the invention disclosed herein without departing from the scope and spirit of the invention. For example, those skilled in the art will recognize that the invention may suitably be practiced using a variety of different bacteria, bacteriophage, and sequencing methods within the general descriptions provided.

The invention illustratively described herein suitably may be practiced in the absence of any element or elements, limitation or limitations which is not specifically disclosed herein. Thus, for example, in each instance herein any of the terms "comprising," "consisting essentially of" and "consisting of" may be replaced with either of the other two terms. The terms and expressions which have been employed are used as terms of description and not of limitation, and there is not intention that in the use of such terms and expressions of excluding any equivalents of the features shown and described or portions thereof, but it is recognized that various modifications are possible within the scope of the invention claimed. Thus, it should be understood that although the present invention has been specifically disclosed by preferred embodiments and optional features, modification and variation of the concepts herein disclosed may be resorted to by those skilled in the art, and that such modifications and variations are considered to be within the scope of this invention as defined by the appended claims.

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In addition, where features or aspects of the invention are described in terms of Markush groups or other grouping of alternatives, those skilled in the art will recognize that the invention is also thereby described in terms of any individual member or subgroup of members of the Markush group or other group. For example, if there are alternatives A, B, and C, all of the following possibilities are included: A separately, B separately, C separately, A and B, A and C, B and C, and A and B and C. Thus, for example, for the bacteria and phage specified herein, the embodiments expressly include any subset or subgroup of those bacteria and/or phage. While each such subset or subgroup could be listed separately, for the sake of brevity, such a listing is replaced by the present description.

Thus, additional embodiments are within the scope of the invention and within the following claims.

## Table 1

# Phages against human and animal pathogenic bacteria

I. Pathogen name	Phage name	II.	Cat alo g#	Origin/reference
Acinetobacter calcoaceticus	A3/2 A10/45 A36 B9GP B9PP BS46 E13 E14 531 Ap3		<b>5</b> "	Felix d'Herelle Reference Centre, Quebec, Quebec  J. Bacteriol 1984, 157: 179-183
Acinetobacter haemolyticus	P78			J. Gen. Microbiol 1986.132: 2633-2636  Felix d'Herelle Reference Centre,Quebec,Quebec
Acinetobacter johnsonii				Felix d'Herelle Reference Centre.Quebec,Quebec
Acinetobacter sp.	BP1 G4, HP2, HP3 & HP4 A1, A4, A9 &			J.Virol.1968.2:716-722 Can.J.Microbiol.1966.12:1023-1030 & J.Virol.1974.13:46-52 & Arch.Virol.1994.135:345-354 Arch.Virol.1994.135:345-354
	196 HP1 A19, A23, A29, A31, A33, A34, A3759 & 2845			Can.J.Microbiol.1966.12:1023-1030  J.Microsc (Paris) 1973.16:215-224 &  CR.Hebdo Seances Acad.Sci.Ser D.Sci Natur(Paris)278:1907-1909 &  Arch.Virol.1994.135:345-354 &  Rev.Can.Biol.1970.29:317-320
Actinobacillus actinomycetecomitans				FEMS Microbiol Lett 1994. 119:329=337

Infec. Immun. 1982. 35: 343-349				/
Actinomyces viscosus  Actinomyces viscosus  Actinomyces viscosus  Actinomyces viscosus  Infect.Immun.1985.48:228-233  Infect.Immun.1988.56:54-59  Infect.Immun.1988.56:54-59  Plasmid 1997.37:141-153  Aeromonas hydrophila  Aeromonas hydrophila  Aeromonas hydrophila  Aeromonas hydrophila  Aeromonas hydrophila  Aeromonas hydrophila  PM2** & PM3  Aeromonas hydrophila  Aeromonas hydrophila  Aeromonas hydrophila  Aeromonas hydrophila  PM2** & PM3  Aeromonas hydrophila  Aeromonas hydrophila  Aeromonas hydrophila  PM4  PM5  PM6				Infec. Immun. 1982. 35: 343-349
Actinomyces viscosus  Actinomyces viscosus  Actinomyces viscosus  Actinomyces viscosus  Infect.Immun.1985.48:228-233  Infect.Immun.1988.56:54-59  Infect.Immun.1988.56:54-59  Plasmid 1997.37:141-153  Aeromonas hydrophila  Aeromonas hydrophila  Aeromonas hydrophila  Aeromonas hydrophila  Aeromonas hydrophila  Aeromonas hydrophila  PM2** & PM3  Aeromonas hydrophila  Aeromonas hydrophila  Aeromonas hydrophila  Aeromonas hydrophila  PM2** & PM3  Aeromonas hydrophila  Aeromonas hydrophila  Aeromonas hydrophila  PM4  PM5  PM6				
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Aeh1 Aeh2 PM5 PM6  Felix d'Herelle Reference Centre,Quebec,Quebec	Aeromonas hydrophila	PM2** & PM3		FEMS Microbiol.Lett. 1990.57:277-282
Ach2   Centre,Quebec   PM4   PM5   PM6   PM6				
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Aeromonas	3		Felix d'Herelle Reference
salmonicida	25		Centre, Quebec, Quebec
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	55R.1		Can. J. Microbiol. 1983. 29: 1458-1461
Alteromonas espejiana	PM2**	27025-B1	The American Type Culture Collection
Asticacaulis			Felix d'Herelle Reference
biprosthecum			Centre,Quebec,Quebec
Asticcacaulis		15261-B1	The American Type Culture Collection
excentricus		15261-B2	
		15261-B3	
	фАс21		
	φΑς24		
Azotobacter vinelandii	ΨΑ024	12518-B1	The American Trans Culture College
Azotobacier vinetanati			The American Type Culture Collection
		12518-B4	
	A14	12518-B5	
	A21	12518-B9	
	A31	12518-B10 13705-B1	
	A41	13/03-61	
	PAVI		
Azotobacter sp.			Virology 1972.49:439-452
			, <b>g,</b>
Bacteroides fragilis	Bf-1		Rev. Infect. Dis. 1979. 1: 325-336
	B40-8		FEMS Microbiol. Lett. 1991. 66: 61-67
	HSP40		Appl. Environ. Microbiol. 1989. 55: 2696- 2701
	phiA1		Zentralbl.bakteriol.1972.222:57-63
Bdellovibrio bacteriovorus	MAC-1		J. Gen. Microbiol. 1987. 133: 3065-3070
Bdellovibrio sp.	VL-1		J.Virol.1973.12:1522-1533
Bordetella	214		Zh.Mikrobiol.Epidemiol.Immuno. 1987.5:9-
brochiseptica			13

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Bordetella			Felix d'Herelle Reference
parapertussis			Centre, Quebec, Quebec
			Mol. Gen. Mikrobiol. Virusol. 1988.4: 22-25
•			Zh.Mikrobiol.Epidemiol.Immuno. 1987.5:9-
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	41405		Zh.Mikrobiol.Epidemiol.Immuno. 1987.5:9-
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Brucella abortus			Felix d'Herelle Reference
			Centre, Quebec, Quebec
		23448-B1	The American Type Culture Collection
		23448-B2	ļ
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	BK-2, TB &		Zh.Mikrobiol.Epidemiol.Immunobiol.1983.2:
	Fi**		48-52
	R/c & R/O		Dev. Biol. Stand. 1984.56: 55-62
Brucella canis	R/c		Dev. Biol. Stand. 1984.56: 55-62
		23456-B1	The American Type Culture Collection
Brucella melitensis	BK-2	23430-D1	
Brucella suis	Wb		Zentralbl.Veterinarmed.1975.22:866-867
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	Fi** & TB		Zh.Mikrobiol.Epidemiol.Immunobiol.1983.2: 48-52
Brucelia sp.			Can. J. Vet. Res. 1989.53: 319-325
			Res. Vet. Sci. 1988. 44: 45-49
	R		Zh.Mikrtobiol.Epidemiol.Immunobiol.1983.2:
Campylobacter coli		43133-B1	The American Type Culture Collection
		43134-B1	
Campylobacter coli	18	43135-B1	The American Type Culture Collection
(Cont'd)	19 20	43136-B1	
Campylobacter jejuni	1	35918-B1	The American Type Culture Collection
Campy Country July	2	35919-B1	
	3	35920-B1	
	4	35921-B1	
	5	35918-B2	
	6	35920-B2	
	7	35922-B2	
	8	35923-B1	
	9	35924-B1	
	10	35925-B1	
	111	35925-B2	
	12	35922-B2	
	13	35924-B2	
	14	35922-B3	
	17	43133-B1	
	18	43134-B1	
	19	43135-B1	
	20	43136-B1	
Campylobacter	HP1		J. Med. Microbiol.1993. 38: 245-249
(Helicobacter) pylori	Ch-1**	-	J. Gen. Virol. 1989. 70: 3381-3390
Chlamydia psittaci	Chp1**		J.Bacteriol.1993.175:3838-3843
Clostridium acetobutylicum	CAK-1		J.Dacteriol. 1773. 173:3030-3043

Clostridium botulinum	1	<del> </del>	Nucleic Acids Res.1990.18:1291
Clostridium botulinum			Nucleic Acids Res. 1990.18:1291
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			Bioch.Biophys.res.Commun.1990.171.1304-
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Clostridium difficile	41 & 56		I Clini Microbial 1095 21:251 254
Сюзігіант аўўсіне	41 & 30	L.,	J. Clini.Microbiol. 1985.21:251-254

Clostridium			Rev.Can.Biol.1977.36:205-215
perfringens			
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			FEMS Microbiol.Lett. 1990.54:323-326
Clostridium		8074-B1	The American Type Culture Collection
sporogenes	59	17886-B1	
ļ	70	17886-B3	
	71	17886-B4	
	72S	17886-B5	
	72L	17886-B6	
Clostridium tetani	A&B		Rev.Can.Biol.1978.37:43-46
Corynebacterium		i	Vopr.Virusol.1986.31:577-584
diphteriae			
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Corynebacterium	NN	12319-B1	The American Type Culture Collection
pseudotuberculosis			
Corynebacterium sp	DLC 2921/49	12052-B1	The American Type Culture Collection

Enterococcus faecalis	42	19948-B1	The American Type Culture Collection
Enterococcus faecium	124	19950-B1 19953-b2 19953-B1	The American Type Culture Collection
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Escherichia coli		11303-B14	The American Type Culture Collection
		11303-B10	
		11303-B21	
		8677-B1	•
		11303-B13	
		13706-B4	
Escherichia coli		15766-B1	The American Type Culture Collection
(Cont'd)		15766-B1	
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	E1	15669-B1	
	u**	15597-B1	
	f2**	21816-B1	
	FCZ	23724-B9	
	fd**	15593-B1	
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		29746-B1	· ·
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		11303-B35	
		11303-B34	
	MS2**	11303-B36	
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	Mu-1	13706-B5	
	Ox6	11303-B1	
	P1**	11303-B2	
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	Q-β**	11303-B4	
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	ZJ/2	35060-B3	
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147 JV1 JV47 JV375 z3**  ** LC-17 LSUS P-3 LSUS J-6 LSUS J-6 LSUS A-11 Lind p92 pR dV-1 dX174** dXcs70am-3 G4** & dK** BF23**	11303-B17 11303-B15 11303-B11 11303-B18 13706-B2 23724-B2 23724-B3 23724-B4 23724-B5 23724-B6 23724-B6 23724-B7 23724-B8 35860-B1 13706-B3 15597-B2 13706-B1 49696-B1	Biochim.Biophysica Acta.1992.1130:277-288  J.Bacteriol.1977.129:265-275
JV1 JV47 JV375 x3**  ** LC-17 L sus P-3 L sus R-5 L sus J-6 L sus A-11 L ind  692  RR 6V-1 6X174** 6Xcs70am-3 G4** & 6K**	11303-B11 11303-B18 13706-B2 23724-B2 23724-B1 23724-B3 23724-B4 23724-B5 23724-B6 23724-B7 23724-B8 35860-B1 13706-B3 15597-B2 13706-B1	<del></del>
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		J.Mol.Biol.1991.218:705-721
		FEBS Lett.1987.215:145-150
Rb18**, Rb51 &		J.Bacteriol.1990.172:180-186
11**, H3, H8, (9,		Mol.Gen.Genet.1990.221:491-494
(18 & Ox1		
M1**, TuIa** & TuIb**		J.Mol.Biol.1987.196:165-174
C10		J.Bacteriol.1979.140:680-686
Qsr'		J.Bacteriol.1985.162:256-262
3278		J.Gen.Microbiol.1988.134:1333-1338
ohi 80**		FEMS Microbiol.Lett.1994.119:71-76
ohi m173		Genetika 1985.21:673-675
f-1		J.Gen.Microbiol.1987.133:953-960
4 & phiR73		Mol.Microbiol.1995.18:201-208
2-2	.=-	J.Gen.Microbiol.1982.128:2797-2804
PRD1		Virology 1990.177:445-451
C3hx		Mol.Gen.Genet.1987.206:110-115
33J**&		Infect.Immunity.1986.53:135-140
H19-B**		J.Bacteriol.1987.169:4308-4312
Ccp-111		Zentralbnl.Bakteriol.Mikrobiol.Hyg,1988.270: 41-51
H   C   C   C   C   C   C   C   C   C	b69** 1**, H3, H8, 9, 18 & Oxl (1**, TuIa** & ulb** 10 sr' 278 ni 80** ni m173 -1 4 & phiR73 -2 RD1 3hx 33J**& 33W**	p17 3** & Ox2** b18**, Rb51 & b69**  1**, H3, H8, 9, 18 & Ox1 (1**, TuIa** & uIb** 10 sr' 278 ni 80** ni m173 -1 4 & phiR73 -2 RD1 3hx 33J**& 33W** 19-B**

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	N4**	Vet.Microbiol.1992.30:203-212
	Phi 80 trp	Ann.Inst.Pasteur.1971.120:121-125
	Obeta 1	J.Bacteriol.1978.133:172-177
	P1CM	J.Gen.Microbiol.1978.107:73-83
	PA-2**	J.Bacteriol.1990.172:1660-1662
	186**	Mol.Gen.Genet.1982.187:87-95
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	21**	Virology 1983.129:484-489
	P4**	MicrobiolRev.1993.57:683-702
	82**	J.Biol.Chem.1987.262:11721-11725
	PSP3	J.Bacteriol.1996.178:5668-5675
*	HK022**	Nucleic Acids Res.1994.22:354-356
		Nucleic Acids Res.1994.22.334-330  Nucleic Acids Res.1986.14:3813-3825
	D108**	· · · · · · · · · · · · · · · · · · ·
Escherichia coli	Rb49	J.Mol.Biol.1997.267:237-249
(Cont'd)	Ike**	J.Mol.Biol.1985.181:27-39
	P22dis	Mol.Gen.Genet.1978.166:233-243
	N15**	J.Bacteriol.1996.178:1484-1486
	If1**	Proc.R.Soc.Lond.B.Biol.Sci.1991.245:23-30
	Stx2Phi-I &	Infect.Immun.1998.66:4100-4107
	Stx2Phi-II	111 1 1000 156 100 106
	18	Virology 1987.156:122-126
	X AC3	J.Gen.Microbiol.1981.126:389-396  Mol.Microbiol.1991.5:715-725

	BW-1 C-1 E920g Esc-7-11 H19J Haiti		Felix d'Herelle Reference Centre,Quebec,Quebec
	HK243 Iα K20 K30 KL <sub>3</sub> M Mu**		·
	O103 O157:H7 P1D pt1 PilHα PR64FS PR772		
	SS4 β4Q λvir** Ω8 09-1 92		
Haemophilus	HP1**		Nucleic Acids Res. 1996.24:2360-2368
influenzae	S2**		Gene 1997. 196: 139-144
Halobacterium cutirubrum	S45	<u>.</u>	Felix d'Herelle Reference Centre,Quebec,Quebec
Halobacterium halobium			Felix d'Herelle Reference Centre,Quebec,Quebec
			Can.J.Microbiol.1982.28:916-921
Halobacterium salinarium			Biol.Chem.Hoppe Seyler 1994.375:747-757

Klebsiella oxytoca	tf-1		J.Gen.Microbiol.1987.133:953-960
Klebsiella pneumoniae	60 92	23356-B1 23357-B1	The American Type Culture Collection
	K19Q		Felix d'Herelle Reference Centre, Quebec, Quebec
	FC3-1 & FC3-9		Can.J.Microbiol.1991.37:270-275
	FC3-10		FEMS Microbiol.Lett.1991.67:291-297
Klebsiella sp.	K11**	· · · · · · · · · · · · · · · · · · ·	Mol.Gen.Genet. 1990.221:283-286
Leptospira sp.	LE1, LE3 & LE4		Res.Microbiol.1990.141:1131-1138
Listeria	243	23074-B1	The American Type Culture Collection
monocytogenes	197,1313 & 9425		Appl.Environ.Microbiol.1997.63:3374-3377
	H387 & H387-A		Appl.Environ.Microbiol.1993.59:2914-2917
	5775,6223 &12682		APMIS.1993.101:160-167
	2389, 2671,		Intervirology 1994.37:31-35 &
	4211 & 2685		Zentralbl.Bakteriol.Mikrobiol.Hyg.1986.261:1 2-28
	4b, 4ab, 4g & 3c		Ann.Microbiol (Paris) 1977.128:185-198
	A118, A500 & A511**		Mol.Microbiol. 1995.16:1231-1241-992
	1, 3, 4, 5, 6, 7, 8, 9, 10, 11, 14, 15, 16, 17, 19 & 20		Ann.Microbiol. (Paris) 1979.130B:179-189
	1/2a, 1/2b, 3c, 4ab, 6a & 6b		Clin.Invest.Med.1984.7:229-232
	φLMUP35 2685		Felix d'Herelle Reference Centre, Quebec, Quebec
Listeria innocua	4211		Felix d'Herelle Reference Centre, Quebec, Quebec
Micrococcus luteus		4698-B1 4698-B4	The American Type Culture Collection
	N3	4698-2	
	N4 N8	4698-B3	
Micrococcus luteus	N17		Can.J.Microbiol. 1979.25:1027-1035
Mycobacterium	BK-3	27203-B1	The American Type Culture Collection
smegmatis	Bol**	27204-B1	
J	Bo 6	27205-B1	and the state of t
	Bo 6II	27205-B2	
	Bo 6III	27205-B3	
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	Mc-4	607-B7	
	NN	11727-B1	
	Phagus lacticola	11759-B1	
	R1	607-B1	

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			Mol.Microbiol.1993.7:395-405
			J.Mol.Biol.1998.279:143-164
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			Proc.Natl.Acad.Sci USA.1988.84:2833-2837
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	29M, 31M, 122,		Arch.Virol.1993.133:39-49 &
	154, 37, 29D, 46,		
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	74D, AG1 &		
	DS6A		
Mycobacterium		23052-B1	The American Type Culture Collection
fortuitum		27207-B1	
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Mycobacterium leprae	T	T .	Ann.Microbiol. (Paris) 1982.133:93-97
Mycobucierium teprae			Am. Microbiol. (Fails) 1982.133.93-97
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Mycobacterium		25618-B1	The American Type Culture Collection
tuberculosis		25618-B2	,,,
	DS6A	4243-B1	
	110 120 % 220		Arch Virol 1002 122:20 40
	110, 139 & 33D		Arch. Virol. 1993.133:39-49
	AG1,GS4E, BG1,		The Biology of Mycobacteria. Academic Press, Toronto 1982 (Ratledge & Stanford)
	PH & BK1		1982.309-351
Mycobacterium sp	Phagus pellegrini	11760-B1	The American Type Collection Culture
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	B1	23239-B1	- <u></u>
l l	DI	4343 <b>7-D</b> 1	

	TM4, ph60, ph72, PhAE39, phAE40 & Bxb1 C2 18 & I15 63 phlei & butyricum MyF3P-59a Bo2a D4,D28 & D32 HC		Microbiology 1995.141:1173-1181  Experentia 1969.25:1112-1113  J.Gen.Virol.1987.68:949-956  Gruzlica 1968.36:617-622  J.Gen.Virol.1975.29:235-238  Z.Allg.Mikrobiol.1968.8:29-37  J.Gen.Virol.1973.20:75-87  J.Exptl.Med.1966.123:327-340  J.Bacteriol.1963.86:608-609
Mycobacterium vaccae	B5	15483-B1	The American Type Culture Collection
Mycobacterium phlei	NN Bo 2 Bo 2h Bo 3	11728-B1 11758-B1 27086-B2 27086-B1	The American Type Culture Collection
Mycoplasma arthritidis	MAV1**		Infect.Immunity.1995.63:4016-4023
Mycoplasma hyorhinis	Hr-1		Arch.Virol.1983.77:81-85
Mycoplasma pneumoniae	Br-1		Arch.Virol.1983.75:1-15
Mycoplasma pulmonis			Plasmid 1995. 33: 41-49
Mycoplasma sp.			J.Gen.Microbiol.1985:131:3117-3126
			J. Virol.1986.59:584-590
			Gene 1994. 141: 1-8

	Microbios 1990. 64: 111-125
	Infection& Immunity 1995. 63: 4016-4023
	Med.Biol.1982.60:116-120
MV-L2 &	Arch.Virol.1979.61:289-296
	Acta.Virol.1978.22:443-450
	J.Gen.Virol.1979.42:315-322
	Virology 1973.55:118-126

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			Science 1971.173:725-727
	,		<del>-</del>
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			J.Clin.Microbiol.1976. 4:87-91
Neisseria perflava			7.CIII.,WICIOSIOI.1370. 4.07-31
Nocardia erythrypolis	φС		J.Gen.Virol.1974.23:247-254
	φEC		J.Bacteriol.1976.126:1104-1107
Pasteurella multocida	B225		Arch.Exp.Veterinarmed.1981.35:433-436
	B939a		Am.J.Vet.Res.1978.39:1565-1566
	Nos.115, 32, 967		Vet.Med.Nauki. 1977.14:33-36
	&		
	1075	20000 71	The American College of College
Propionibacterium	NN	29399-B1	The American Type Collection Culture
acnes			

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Pseudomonas		12175-B1	The American Type Culture Collection
aeruginosa	2	12175-B2	
	2A	12175-B3	
	2B	12175-B4	
	11	14205-B1	
	16	14206-B1	
	24	14207-B1	
	27	14208-B1	
	44	14209-B1	
	73	14210-B1	
	95	14211-B1	
	109	14212-B1	
	113	14213-B1	
	249	14214-B1	
	B3	15692-B1	
	Hoff 2	14203-B1	
	Hoff 3	14204-B1	
	Pa	12055-B1	
	Pb	12055-B2	
	PB-1	15692-B3	
	Pc	12055-B3	,
	Pf	25102-B1	
	PP7**	15692-B2	
	417		Felix d'Herelle Reference
			Centre, Quebec, Quebec
	ł		,
	7 0 21		
	7 & 31	1	
	Pf3**		J.Virol.1983.47:221-223
	ф-МС		Can.J.Microbiol.1969.15:1179-1186
	Pf1**		J.Mol.Biol.1991.218:349-364
	PR4**		J.Gen.Virol.1979.43:583-592
	A7		J.Bacteriol.1992.174:2407-2411
	KF1		J.Biochem. 1983.93:61-71
	¢CTX**		Mol.Microbiol.1993.4:1703-1709
			J.Virol.1977.24:135-141
	f2**		J.VIG1.19/7.24:135-141
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 φKZ, 21, φNZ, PMN17, PTB80,	dd
68, PB-1, E79,	
16,	
109, 352, 1214,	
F8, 71, 337, M4,	
φC17, SL2, B17,	
Li-24, φmnP78,	
PS17**, φ1, 73,	
M6, Li-2, 7,	
φmnF82,	
PTB2, PTB20,	
PTB42, φKF77,	
31, PTB21, 119x,	
φPLS27, B3,	
258,	
Hw12, PM57,	
PM62, PM105,	
148, PM681,	
198,	
218, 222, 242, 246,	
PC131, φC11,	
SL5,	
D3112**, Jb19,	
F7,	
PM69, PM13,	
PM61, PM113,	
φ240, 249 & 269	

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Pseudomonas aeruginosa	297, 309, 318, 11,	Arch.Virol.1993.131:141-151
(Cont'd)		
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Pseudomonas cepacia			Felix d'Herelle Reference
			Centre, Quebec, Quebec
	i		
Pseudomonas fragi		27362-B1	The American Type Culture Collection
1 scanomonas ji ug.		27363 B1	The Interiorn Type Canala Concensis
	wy		
Pseudomonas	ф6	1	Felix d'Herelle Reference
phaseolicola			Centre, Quebec, Quebec
Pseudomonas putida	gh-1	12633-B1	The American Type Culture Collection
Pseudomonas syringae		40492-B1	The American Type Culture Collection
	1.0	21781-B1	
p	φ-6	40790 D1	The American Trans Culture City
Pseudomonas sp.	PPs-G3 Sab 2	49780-B1	The American Type Culture Collection Felix d'Herelle Reference
Salmonella bareilly	Sab 2		Centre, Ouebec, Ouebec
Salmonella enteritidis	1, 2,3 & 6	1	Epidemiol.Infect.1995.114:227-236
	2a, 3a, 4a, 5a, 6a,		Vet.Med.Nauki.1975.12:55-60
	7a, 8a, 9a, 15,		1 0011200.110010112.03 00
···	19, 20 &21**		
Salmonella newington	Epsilon 34		J.Struct.Biol. 1995.115:283-289
Salmonella newport		27869-B1	The American Type Culture Collection
		27869-B2	
	16-19		}
			Felix d'Herelle Reference
	i		Centre, Quebec, Quebec
Salmonella paratyphi		19940-B1	The American Type Culture Collection
	Paratyphoid A	12176-B1	
	Jersey		Felix d'Herelle Reference
	Jersey		Centre, Quebec, Quebec
Salmonella	SasL1, SaL2, Sal		Indian J.Med.Res. 1997.105:47-52
senftenberg	3,		
	SaL4, SaL5 &		
	SasL6		
Salmonella	P22**	19585-B1	The American Type Culture Collection
typhimurium	SL-1	40282	
	MB78**		J.Virol. 1982.41: 1038-1043
	SE1		J.Gen.Microbiol.1986.132:1035-1 <del>0</del> 41
	LT2		Virology 1971.45:835-636
		l	
	ES18**		Virology 1970.42:621-632 J.Virol.1985.56:1034-1036

	P1CM cir-100		Mol.Gen.Genet.1975.138:113-126
-0.0	F22	<del> </del>	Genet.Res.1986.48:139-143
	Fels 1		J.Gen. Virol. 1978.38:263-272
	Fels 2		Genet.Res.1986.48:139-143
	Px	1	Mol.Gen.Genet.1970.108:184-202
	Plkc	<del> </del>	
	A3 & A4	<del> </del>	Virology 1974.60:503-514
	<del></del>	<del>                                     </del>	J.Bacteriol. 1987.169:1003-1009
	HT		Genet.Res.1976.27:315-322
Salmonella	IRA		J.Basic Microbiol. 1990.30:707-716
typhimurium	Mudl		Mol.Gen.Genet. 1986.202:327-330
(Cont'd)	P22 (cir4-1, cir5- 1 & cir6-1)		Mol.Gen.Genet.1984.198:105-109
	BF23**		Mol.Gen.Genet.1976.147:195-202
	Kbl		J.Bacteriol.1974.117:907-908
	P221dis		J.Gen.Virol.1978.41:367-376
	PRD1**		Virology 1990.177:445-451
	I <sub>2</sub> -2**		J.Gen.Microbiol.1982.128:2797-2804
	tf-1		J.Gen.Microbiol.1987.133:953-960
	X**		J.Gen.Microbiol.1981.126:389-396
Salmonella	8	19937-B1	The American Type Culture Collection
typhosa/typhi	23	19938-B1	,,,
	25	19939-B1	
	46	19942-B1	
	53	19943-B1	
•	163	19946-B1	
	175	19947-B1	
	ViI	27870-B1	
	ViVI	27870-B2	
	01		Felix d'Herelle Refrence Centre, Quebec, Quebec
	ViII		Chung Hua Liu Hsing Ping
			H.T.C.1992.13:288
	j2		J.Gen.Microbiol.1983.129:3395-33400
Salmonella sp.	P3	25957-B1	The American Type Culture Collection
	P4**	25957-B2	
	P9a	25957-B3	
	P9c	25957-B4	
	P10	259 <b>5</b> 7-B5	
	102	19945-B1	
	Chi (χ)	9842-B1	
	R34	97541	
	MG40		Virology 1968.34:521-530
	P14		Microb.Pathog.1990.8:393-402
	PSP3		Virology 1992.188:414
	Ike**		Zentralbl.Bakteriol.1976.234:294-304
·	P27 & 9NA		J.Virol.1986.12:921-931
Sphaerotilus natans	SN1		Appl.Environ.Microbiol.1979.37:1025-1030

Shigella dysenteriae		23351-B1	The American Type Culture Collection
	P2	11456b	
	<i>\$</i> -80	11456a-B1	
Shigella flexeneri	D20	12661-B1	The American Type Culture Collection
	SfII**		Mol.Microbiol.1997.26:939-950
	SfV**		Gene 1997.22:217-227
	Sf6**		Mol.Microbiol.1995.18:201-208
	SfX	<u> </u>	Gene 1993.129:99-101
Shigella sonnei	C16**		
	Ufa		MolBiol (Mosk) 1977.11:323-331
Shigella sp	37	23354-B1	The American Type Culture Collection
Spiroplasma citri	SpV1		Plasmid 1993.29:193-205
Spiroplasma sp.	SpV1-R8A2B		Nucleic Acids Res. 1990.18:1293
	SpV3		Isr.J.Med.Sci.1987.23:429-433
	Sp V4		J.Bacteriol.1987.169:4950-4961
Staphylococcus albus			Staphylococci & Staphylococcal Infections.1997.
			Vol1:503-508 (Karger, Basel)
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Staphylococcus aureus		27702-B1 27703-B1	The American Type Culture Collection
		27704-B1	
		23360-B1	
	1.5	23361-B1	
	15	27705-B1	
	17	27712-B1	
	29	27690-B1	
	42D**	27691-B1	
	42E	27692-B1	
	47	27693-B1	
	52	27694-B1	
	52A	27695-B1	
	53	27696-B1	
	54	27697-B1	
	55	27698-B1	
	71	27699-B1	
	75	27693-B2	
	77	27700-B1	
	79	27701-B1	
	80	27706-B1	
	81	27707-B1	
	83A	27708-B1	
	84	33742	
	85**	33741-B1	
	88	15565	
	92	19685-B1	
	5504'	11987-B1	
	K	.11988-B1	
	P1	15752-B1	
	P14	Í	
	UC18	Ì	

	Twort**	HER 101 HER 239 HER 283 HER 49	Felix d'Herelle Reference Centre,Quebec,Quebec
	φ11**		J.Bacteriol.1988.170:2409-2411
	ф13** & ф42**		J.GenMicrobiol.1989.135:1679-1697
	L54a**		J.Bcteriol.1986.166:385-391
	80α**		Can.J.Microbiol.1996.43:612-616
	94,95 & 96		J.Clin.Microbiol.1988.26:2395-2401
	φ131,A <sub>3</sub> & A <sub>5</sub>		Staphylococci & Staphylococcal Infections.1997.
	DI: DI // ++		Vol1:503-508 (Karger,Basel)
Cambulana	Phi PVL**		Gene 1998.215:57-67 Felix d'Herelle Reference
Staphylococcus carnosus	BaSTC2		Centre, Quebec, Quebec
Staphylococcus epidermidis	1a, 2b, 3a, 4b, 5a, 6b, 7b, 8c, 9a, 10a, 11b,12a & 13b		Can.J.Microbiol.1988.34:1358-1361
	41, 63, 118II, 138, 245, 336, 392 & 550		Res.Virol.1994.145:111-121
Staphylococcus	1154A, 1405,		Res.Virol.1990.141: 625-635 &
saprophyticus	1314, 1139 & 1259		Res.Virol.1994.145:111-121
Staphylococcus sp.	Phi 812, Phi 131, SK311 & U16		Virology 1998.246:241-252
Streptococcus faecalis	VD13	HER44	Felix d'Hereile Reference Centre, Quebec, Quebec
Streptococcus faecium	PE1		Zentralbl.Bakteriol.1975.231:421-425
Streptococcus oralis	Cp-1** & Cp- 7**		FEMS Microbiol.Lett.1989.65:187-192

Streptococcus pneumoniae	Cp-1**	HER223	Felix d'Herelle Reference Centre,Quebec,Quebec
	C- 1** C- 5**	<del></del>	
	Cp-1**, Cp-5**, Cp-7**, Cp-9**,		J.Virol.1981.40:551-559 &
	ω-1 & ω-2		Eur.J.Biochem.1979.101:59-64 &
			Microbial Drug Resistance 1997.3:165-176
	HB-623 & HB- 746		J.Virol.1990.64:5149-5155
	EJ -1**		J.Bacteriol.1992.174:5516-5525
	Dp-2 & Dp-4		J.Virol.1978.26:221-225
	Dp-1		Virology 1975.63:577-582
	ω-3 & ω-8		J.Virol.1976.19:659-667
•	304		J.Bacteriol.1980.141:1298-1304
	HB-1,HB-2,		J.Bacteriol.1979.138:618-624
	HB-3**,		
	HB-4, HB-5 &		
	HB-6	l	
Streptococcus	T12**		Mol. Microbiology. 1997#23:719-728
pyogenes	A-1	12202-B1	The American Type Culture Collection
	A-6	12203-B1	,
	A-25	12204-B1	
	Kjem	14918	
Streptococcus	1	HER 339	Felix d'Herelle Refrence
sp./Enterococcus	182	HER 80	Centre, Quebec, Quebec
	VD1884	HER 323	
	1A	12169-B1	The American Type Culture Collection
	1B	12170-B1	
	NN	21597-B1	
	42	19948-B1	
	118	19951-B2	
	120	19952-B1	
Veillonella rodentium	N2		Antonie Van Leeuwenhoek 1989.56:263-27
Vibrio cholerae	Psi 92		Intervirology 1993.36:237-244
	VCB-1,2,3 & 4		J.Infetion 1998.36:131
	CP-Ti**		J.Virol.1984.51:163-169
	VSK		FEMS Microbiol.Lett.1996.145:17-22
	Phi138		J.Virol.1986.57:960-967
	Phi149		J.Virol.1985.140:217-223
	Fs-2**		Microbiology 1998.144:1901-1906
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138	14100-B1	The American Type Culture Collection
145	14100-B2	
149	14100-B30	
163	14100-B4	
N-4	51352-B1	
S-5	51352-B2	
S-20	51352-B3	
M-4	51352-B4	
D-10	51352-B5	
1	51352-b6	
l II	51352-B7	
ш	51352-B8	
lıv	51352-B9	
v	51352-B10	
UTAK		Felix d'Herelle Reference
		Centre, Quebec, Quebec
e4		J.Gen.Virol.1987.68:1411-1416
nt1,nt6		Felix d'Herelle Reference
		Centre, Quebec, Quebec
KVP40**		Felix d'Herelle Reference
VF33		Centre,Quebec,Quebec
VP1		
φ60		
φHAWI-5		
φPEL8C-1		
α3a		Felix d'Herelle Reference Centre, Quebec, Quebec
NN	11985-B1	The American Type Culture Collection
ph1	51582-B1	
I PIXX		
Phi149		J.Virol.1987.61:3999-4006
	K 13 14 16 24 32 57 138 145 149 163 N-4 S-5 S-20 M-4 D-10 I II III IV V UTAK  e <sub>4</sub> nt1,nt6  KVP40** VF33 VP1 φ60 φHAWI-5 φPEL8C-1 α3a	e5 X29 β κ 13 14 16 24 32 57 138 14100-B1 145 14100-B2 149 14100-B30 163 14100-B4 N-4 51352-B1 S-5 51352-B2 S-20 51352-B3 M-4 D-10 51352-B5 I 51352-B6 II 51352-B6 II 51352-B7 III 51352-B8 IV 51352-B9 V 51352-B9 V 51352-B10 UTAK  e4 nt1,nt6  KVP40** VF33 VP1 φ60 φHAWI-5 φPEL8C-1 α3a NN 11985-B1

Yersinia enterocolitica	l i		Felix d'Herelle Reference
Tersinia enteroconica	2		Centre, Quebec, Quebec
	3		
	4		
	5		
	6	-	
	7		
	8		
	9	1	
	φYeO3-12		
	I, IV & VIII		Zentralbl.Bakteriol.Mikrobiol.Hyg.1982.253:1
Yersinia pestis	R	23208-B1	The American Type Culture Collection
, , , , , , , , , , , , , , , , , , ,	s	11593-B1	
	Y	23053-B1	
	II		Zh.Mikrobiol.Epidemiol.Immunobiol.1990.11
Yersinia pseudotuberculosis	PST**	23207-B1	The American Type Culture Collection
Yersinia sp.	RD2		Mol.Gen.Mikrobiol.Virusol.1990.8:18-21

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Table 2

>Bacteriophage 77, complete genome sequence, 41708 nucleotides

```
gatcaaaata cttggggaac ggttagggag taaacttcgc gataatttta aaaattcatg
      tataaccccc ctcttataac cattttaagg caggtgatga aatggagatt atagtcgatg
61
      aaaatttagt gcttaaagaa aaagaaaggc tacaagtatt atataaagac atacctagca
121
      ataaattaaa agtagttgat ggtttaatta ttcaagcagc aaggctacgt gtaatgcttg
181
      attacatgtg ggaagacata aaagaaaaag gtgattatga tttatttact caatctgaaa
241
      aggcgccacc atatgaaagg gaaagaccag tagccaaact atttaatgct agagatgctg
301
      catatcaaaa aataatcaaa caattatcgg atttattgcc cgaagagaaa gaagacacag
361
      aaacgccatc tgatgattac ctatgattag taataaatac gttgatgaat atataaattt
421
      gtggaaacaa ggaaagataa ttttaaataa agaaagaatt gatctcttta attatctaca
481
      aaaacatata tattcacgag atgatgtata ttttgatgaa cagaaaatcg aggattgtat
541
      caaatttatt gaaaaatggt attttccaac attaccattt caaaggttta tcatagctaa
601
      tatatttett atagataaaa atacagatga agetttettt acagaatttg etatttteat
661
      gggacgtgga ggcgggaaaa acggtctaat aagtgctatt agtgattttc tttctacgcc
721
      cttacacgga gttaaagaat atcacatctc cattgttgct aatagtgaag atcaagcaaa
781
      aacatcgttt gatgaaatca gaaccgtttt aatggataac aaacgaaata agacgggtaa
841
      aacgccaaaa gctccttatg aagttagtaa agcaaaaata ataaaccgtg caactaaatc
901
      ggttattcga tataacacat caaacacaaa aaccaaagac ggtggacgtg aggggtgtgt
961
      tatttttgat gaaattcatt atttctttgg tootgaaatg gtaaacgtca aacgtggtgg
1021
      attaggtaaa aagaaaaata gaagaacgtt ttatataagt actgatggtt ttgttagaga
1081
      gggttatatc gatgcaatga agcacaaaat tgcaagtgta ttaagtggca aggttaaaaa
1141
      tagtagattg tttgcttttt attgtaagtt agacgatcca aaagaagttg atgacagaca
1201
      gacgtgggaa aaggcgaacc caatgttaca taaaccgtta tcagaatacg ctaaaacact
1261
      gctaagcacg attgaagaag aatataacga tttaccattc aaccgttcaa ataagcccga
1321
      attcatgact aagcgaatga atttgcctga agttgacctt gaaaaagtaa tagcaccatg
1381
      gaaagaaata ctagcgacta atagagagat accaaattta gataatcaaa tgtgtattgg
1441
      tggtttagac tttgcaaaca ttcgagattt tgcaagtgta gggctattat tccgaaaaaa
1501
      cgatgattac attiggttag gacattcgtt tgtaagacaa gggtttttgg atgatgtcaa
1561
      attagaacct cctattaaag aatgggaaaa aatgggatta ttgaccattg tcgatgatga
1621
      tgtcattgaa attgaatata tagttgattg gtttttaaag gctagagaaa aatatgggct
1681
      tgaaaaagtc atagctgata attatagaac tgatattgta agacgtgcgt ttgaggatgc
1741
      tggcataaaa cttgaagtac ttagaaatcc aaaagcaata catggattac ttgcaccacg
1801
      tatcgataca atgtttgcga aacataacgt aatatatgga gacaatcctt tgatgcgttg
1861
      gtttactaat aatgttgctg taaaaatcaa gccggatgga aataaagagt atatcaaaaa
1921
      agatgaagtc agacgtaaaa cggatggatt catggctttt gttcacgcat tatatagagc
1981
      agacgatata gtagacaaag acatgtctaa agcgcttgat gcattaatga gtatagattt
2041
      ctaatagagg aggtgagaca tgagtattct agaaaagata tttaaaacta ggaaagatat
2101
      aacatatatg cttgatttag atatgataga agatctatca caacaagcgt atgtgaaacg
2161
      tttagcgatt gatagttgta ttgaatttgt tgcgcgaget gtcgctcaaa gtcattttaa
2221
      agtattggaa ggtaatagaa ttcaaaagaa tgatgtttac tacaagttaa atataaaacc
2281
      aaatactgac ttatcaagcg atagtttttg gcaacaagtt atatataaac taatttatga
2341
      taacgaggtt ttaatcgtag taagtgacag caaagaatta cttatcgcag atagctttta
2401
      cagagaagag tacgctttgt atgatgatat attcaaagat gtaacggtta aagattatac
2461
      ttatcaacgt actttcacaa tgcaagaggt catatattta aagtacaaca acaataaagt
2521
      gacacacttt gtagaaagtc tattcgaaga ttacgggaaa atattcggaa gaatgatagg
2581
      tgcacaatta aaaaactatc aaataagagg gattttgaaa tctgcctcta gcgcatatga
2641
      cgaaaagaat atagaaaaat tacaagcgtt cacaaataaa ttattcaata cttttaataa
2701
      aaatcaacta gcaatcgcgc ctttgataga aggttttgat tatgaggaat tatctaatgg
2761
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36241 ctaactttat tttaaaaggg cggaaacaat gaaaatcaaa attgaaaaag aaatgaattt 36301 acctgaactt atccaatggg cttgggataa ccccaagtta tcaggtaata aaagattcta 36361 ttcaaatgat gttgagcgca actgttttgt gacttttcat gttgatagca tcttatgtaa 36421 tgtgactgga tatgtatcaa ttaacgataa atttactgtt caagaggaga tataacaatg 36481 aaaatcaaag ttaaaaaaga aatgagatta gatgaattaa ttaaatgggc gcgagaaaat 36541 coggatotat cacaaggaaa aatatittit toaacaggat tiagtgatgg attogttogt 36601 tttcatccaa atacaaataa gtgttcgacg tcaagtttta ttccaattga tatccccttc 36661 atagttgata ttgaaaaaga agtaacggaa gagactaagg ttgataggtt gattgaatta 36721 ttcgagattc aagaaggaga ctataactct acactatatg agaacactag tataaaagaa 36781 tgtttatatg gcagatgtgt gcctaccaaa gcattctaca tcttaaacga tgacctaact 36841 atgacgttaa totggaaaga tggggagttg otagtatgat gttgaaattt aaagottggg 36901 ataaagataa aaaagttatg agtattattg acgaaatcga ttttaatagt gggtacattt 36961 tgatttcaac aggttataaa agtttcaatg aagtaaaact attacaatac acaggattta 37021 aagatgtgca cggtgtggag atttatgaag gggatattgt tcaagattgt tattcgagag 37081 aagtaagttt tatcgagttt aaagaaggag cettttatat aacttttage aatgtaactg 37141 aattactaag tgaaaatgac gatattattg aaattgttgg aaatattttt gaaaatgaga 37201 tgctattgga ggttatgaga tgacgttcac cttatcagat gaacaatata aaaatctttg 37261 tactaactct aacaagttat tagataaact tcacaaagca ttaaaagatc gtgaagagta 37321 caagaagcaa cgagatgagc ttattgggga tatagcgaag ttacgagatt gtaacaaaga 37381 tctagagaag aaagcaagcg catgggatag gtattgcaag agcgttgaaa aagatttaat 37441 aaacgaattc ggtaacgatg atgaaagagt taaattcgga atggaattaa acaataaaat 37501 ttttatggag gatgacacaa atgaataatc gcgaaaaaat cgaacagtcc gttattagtg 37561 ctagtgcgta taacggtaat gacacagagg ggttgctaaa agagattgag gacgtgtata 37621 agaaagcgca agcgtttgat gaaatacttg agggaatgac aaatgctatt caacattcag 37681 ttaaagaagg tattgaactt gatgaagcag tagggattat ggcaggtcaa gttgtctata 37741 aatatgagga ggaataggaa aatgactaac acattacaag taaaactatt atcaaaaaat 37801 gctagaatgc ccgaacgaaa tcataagacg gatgcaggtt atgacatatt ctcagctgaa 37861 actgtcgtac tcgaaccaca agaaaaagca gtgatcaaaa cagatgtagc tgtgagtata 37921 ccagagggct atgtcggact attaactagt cgtagtggtg taagtagtaa aacgtattta 37981 gtgattgaaa caggcaagat agacgcggga tatcatggca atttagggat taatatcaag 38041 aatgatgaag aacgtgatgg aatacccttt ttatatgatg atatagacgc tgaattagaa 38101 gatggattaa taagcatttt agatataaaa ggtaactatg tacaagatgg aagaggcata 38161 agaagagttt accaaatcaa caaaggcgat aaactagctc aattggttat cgtgcctata 38221 tggacaccgg aactaaagca agtggaggaa ttcgaaagtg tttcagaacg tggagcaaaa 38281 ggcttcggaa gtagcggagt gtaaagacat cttagatcga gttaaggagg ttttggggaa 38341 gtgacgcaat acttagtcac aacattcaaa gattcaacag gacgaccaca tgaacatatt 38401 actgtggcta gagataatca gacgtttaca gttattgagg cagagagtaa agaagaagcg 38461 aaagagaagt acgaggcaca agttaaaaga gatgcagtta ttaaagtggg tcagttgtat 38521 gaaaatataa gggagtgtgg gaaatgacgg atgttaaaat taaaactatt tcaggtggag 38581 tttattttgt aaaaacagct gaaccttttg aaaaatatgt tgaaagaatg acgagtttta 38641 atggttatat ttacgcaagt actataatca agaaaccaac gtatattaaa acagatacga 38701 ttgaatcaat cacacttatt gaggagcatg ggaaatgaat cagctgagaa ttttattaca 38761 tgacggtagt agtttgatat tacatgaaga tgaattattt aacgaaatag tatttgtttt 38821 ggacaatttt agaaatgatg atgactattt aacgatagaa aaagattatg gcagagaact 38881 tgtattgaac aaaggttata tagttgggat caatgttgag gaggcagatg atgattaaca 38941 tacctaaaat gaaattcccg aaaaagtaca ctgaaataat caaaaaatat aaaaataaag 39001 cacctgaaga aaaggctaag attgaagatg attttattaa agaaattaaa gataaagaca 39061 gtgaatttta cagtcctacg atggctaata tgaatgaata tgaattaagg gctatgttaa 39121 gaatgatgcc tagtttaatt gatactggag atgacaatga tgattaaaaa acttaaaaat 39181 atggatgggt tcgacatctt tattgttgga atactgtcat tattcggtat attcgcattg 39241 ctacttgtta tcacattgcc tatctataca gtggctagtt accaacacaa agaattacat 39301 caaggaacta ttacagataa atataacaag agacaagata aagaagacaa gttctatatt 39361 gtattagaca acaaacaaqt cattgaaaat tccgacttat tattcaaaaa gaaatttgat 39421 agcgcagata tacaagctag gttaaaagta ggcgataagg tagaagttaa aacaatcggt 39481 tatagaatac actttttaaa tttatatccg gtcttatacg aagtaaagaa ggtagataaa 39541 caatgattaa acaaatacta agactattat tottactago aatgtatgag ttaggtaagt 39601 atgtaactga gcaagtgtat attatgatga cggctaatga tgatgtagag gcgccgagtg 39661 attacgtctt tcgagcggag gtgagtgaat aatgagaata tttatttatg atttgatcgt 39721 tttgctgttt gctttcttaa tatccatata tattattgat gatggagtga taataaatgc 39781 attaggaatt tttggtatgt ataaaattat agattccttt tcagaaaata ttataaagag 39841 gtagataaaa atgaacgagc aaataatagg aagcatatat actttagcag gaggtgttgt 39901 gctttattca gttaaagaga tttttaggta ttttacagat tctaacttac aacgtaaaaa. 39961 aatcaattta gaacaaatat atccgatata tttagattgt tttaaaaaagg ctaaaaaagat 40021 gattggagct tatattattc caacagaaca gcatgaattt ttagattttt ttgatattga 40081 agtotttaat aatttagata agcaaagtaa aaaagcgtat gaaaatgtta ttggatttag 40141 acaaatgatt aatttatcaa atagagttaa ggcaatggaa gattttaaga tgagtttcaa 40201 caatgaattt agtacaaatc agattttttt taatccttct tttgttatgg aaacaattgc 40261 tattataaat gaatatcaaa aagatatatc ttatttaaaa aatataatta ataaaatgaa

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40321	tgaaaataga	gcttataatc	atattgatag	ttttatcact	tcagagtacc	gacgaaaaat
40381	aaacgattat	aatctttatc	ttgataaatt	tgaagaacag	tttagtcaaa	agtttaaaat
40441	aaacagaact	tcgataaaag	aaagaattat	tattaattta	aacaagagga	gatttaaatg
40501	atgtggatta	ctatgactat	tgtatttgct	atattgctat	tagtttgtat	cagtattaat
40561	agtgatcgtg	caagagagat	acaagcactt	agatatatga	atgattatct	acttgatgaa
40621	gtagttaaaa	ctaaagggta	caacgggtta	gaagaataca	ggattgaatt	gaagcgaatg
40681	aataacgata	ttaaaaagta	atttatatta	tcggaggtat	tgcattgaat	gataaagatt
40741	gagaaacacg	atatcaaaaa	gcttgaagaa	tacattcagc	acatcgataa	ctatcgaaga
40801	gagttgaaga	tgcgagaata	tgaattactt	gaaagtcatg	aaccagataa	tgcgggagct
40861	ggcaaaagta	atttgccggg	taacccgatt	gaacgatgtg	caataaagaa	gtttagtgat
40921	aacaggtaca	atacattaag	aaatatagtt	aacggtgtag	atagattgat	aggtgaaagt
40981	gatgaggata	cgcttgagtt	attaaggttt	agatattggg	attgtcctat	tggttgttat
41041	gaatgggaag	atatagcaca	ttactttggt	acaagtaaga	caagtatatt	acgtagaagg
41101	aatgcactga	tcgataagtt	agcaaagtat	attggttatg	tgtagcggac	ttttacccta
41161	tgtaagtccg	cattaaaaca	gtttattatg	ttagtatcag	attaatattt	aaagttatta
41221	aatgctaata	cgacgcatga	acaagaggcg	catcactatg	tgatgtgtct	ttttatttat
41281	gaggtatgaa	catgttcaaa	ctaattgtaa	atacattact	acacatcaag	tatagatgag
41341	tcttgatact	acttaagtta	tataaggtga	aacattatga	tgactaaaga	cgaacgtata
41401	cgattctata	agtctaaaga	atggcaaata	acaagaaaaa	gagtgctaga	aagagataat
41461	tatgaatgtc	aacaatgtaa	gagagacggc	aagttaacga	catatgacaa	aagcaagcgt
41521	aagtcgttgg	atgtagatca	tatattatcg	ctagaacatc	atccggagtt	tgctcatgac
41581	ttaaacaatt	tagaaacact	gtgtattaaa	tgtcacaaca	aaaaagaaaa	gagatttata
41641	aaaaaagaaa	ataaatggaa	agacgaaaaa	tggtaaatac	ccccgggtca	aaaaaatcaa
41701	aagcgatc					

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152

Table 3

	Name	Position		Name	Position	
1	77ORF005	1957221026	48	77ORF052	17622013	
2	77ORF006	39765196	49	77ORF053	3752137757	
3	77ORF007	2187123076	50	77ORF054	2281823060	
4	77ORF008	21203307	51	77ORF055	1754617788	
5	77ORF009	3194632803	52	77ORF058	1889219122	
6	77ORF010	2609226889	53	77ORF059	3456434785	
7	77ORF011	2444125208	54	77ORF064	2957429795	
8	77ORF012	2978830576	55	77ORF065	2852828746	
9	77ORF012	3362034399	56	77ORF066	2749427703	
10	77ORF014	2776028512	57	77ORF069	3834138547	
11	77ORF014	32914028	58	77ORF070	3626936475	
12			59	77ORF070 77ORF071	4049840701	
	77ORF016	3286733610				
13	77ORF017	2326923982	60	77ORF072	3873538938	
14	77ORF018	3116931840	61	77ORF073	3094531148	
15	77ORF019	3985140501	62	77ORF074	3854438738	
16	77ORF020	69267570	63	77ORF075	1367313870	
17	77ORF021	3776238304	64	77ORF077	2535725605	
18	77ORF022	3060531156	65	77ORF079	2908929280	
19	77ORF023	2690327346	66	77ORF080	3520435389	
20	77ORF024	1070011140	67	77ORF085	2406024242	
21	77ORF025	970710147	68	77ORF092	3970639876	
22	77ORF026	4072941145	69	77ORF094	3222632393	
23	77ORF027	65186925	70	77ORF096	1360613773	
24	77ORF028	3479535199	71	77ORF098	70927256	
25	77ORF029	61176521	72	77ORF102	2905129212	
26	77ORF030	3647836879	73	77ORF104	3439334551	
27	77ORF031	3915139546	74	77ORF109	1828218434	
28	77ORF032	3389234266	75	77ORF112	3954339692	
29	77ORF033	57586120	76	77ORF117	2736127501	
30	77ORF034	78868236	77	77ORF118	3839038530	
31	77ORF035	1925819560	78	77ORF120	3605936199	
32	77ORF036	3687637223	79	77ORF124	3369933833	
33	77ORF037	102446	80	77ORF128	1422114355	
34	77ORF038	3490835219	81	77ORF130	1567515806	
35	77ORF039	3722037528	82	77ORF133	84148542	
36	77ORF040	4137741676	83	77ORF140	1311313235	
37	77ORF041	3545435753	84	77ORF147	70297148	
38	77ORF042	54905774	85	77ORF149	3066830787	
39	77ORF042 77ORF043	2930429564	86	77ORF149 77ORF151	3183731953	
40	77ORF043 77ORF044	1848118768	87	77ORF151 77ORF155	3027830391	
40 41	77ORF044 77ORF045	52165500	87 88	770RF155 770RF157	40444157	
	77ORF045 77ORF046	2566325935				
42 43			89	77ORF167	2069220799	
43	77ORF047	1115911425	90	77ORF175	3571735821	
44 45	77ORF048	2877629039	91	77ORF176	68366940	
45	77ORF049	3601336255	92	77ORF178	3539035491	
46	77ORF050	3575336007	93	77ORF179	83188419	-
47	77ORF051	3893139167	94	77ORF182	2926829564	

47 77ORF051 38931..39167 94 77ORF182 29268..29564

WO 00/32825

153

#### Table 4

## 77ORF017 sequence

2398	2		ato	acg	cat	aat	ata	gaa	aaa	cgc	att	aat	aaa	tta	aaaa	acttct
1	M	T	H	N	I	Ε	K	R	I	N	K	L	K	T	S	
2393	7		gga	aat	cca	aaa	ttt	aaa	aag	tta	gat	tca	gat	att	cact	tattta
16	G	N	P	K	F	K	K	L	D	S	D	I	H	Y	L	
2389	2		ctc	aag	aga	ttt	gaa	ggt	gaa	aaa	aac	cat	aaa	ggt	ttt	tatcca
31	L	K	R	F	E	G	E	K	N	H	K	G	F	Y	P	
2384	7		aag	jttt	aaa	caa	gga	gaa	ata	gtt	ttt	gta	gat	ttc	ggt	ataaac
46	K	F	K	Q	G	E	I	V	F	V	D	F	G	I	N	
2380	2		gtt	aat	aaa	gaa	ttt	tct	aat	tca	cac	ttt	gca	ata	gtga	atgaat
61	V	N	K	E	F	S	N	S	H	F	Α	I	V	M	N	
2375	7		aaa	aat	gat	tct	aat	acg	gag	gat	ata	gta	aat	gtt	att	cctta
76	K	N	D	S	N	T	E	D	I	V	N	V	I	P	L	
2371	2		tcc	tct	aaa	gaa	aac	aaa	aag	tat	tta	aag	atg	aat	ttt	gatttg
91	S	S	K	E	N	K	K	Y	L	K	M	N	F	D	L	
2366	7		aaa	itgg	gag	tat	tat	tta	aga	ttg	ttt	tta	aat	tta	atta	agcgcg
106	K	W	E	Y	Y	L	R	L	F	L	N	L	I	S	Α	
2362	2		caa	aat	aat	tca	gct	ata	tta	aaa	gaa	gtt	ttc	gat	aaaa	aaatac
2362 121	2 Q	N	caa N	aat S	aat A		_				gaa F	_		gat K		aaatac
	Q	N	N	S	A	I	L	K	E	V	F	D	K	K	Y	aaatac attgaa
121	Q 7	N K	N caa	S	A	I	L	K gaa	E ttc	V atc	F	D aaa	K gat	K tat	Y	
121 2357	Q 7 Q		N caa N	S aaa N	A aac T	I aac E	L aca F	K gaa I	E ttc T	V atc K	F act D	D aaa Y	K gat F	K tat I	Y ttta E	
121 2357 136	Q 7 Q		N caa N	S aaa N	A aac T	I aac E gat	L aca F	K gaa I	E ttc T	V atc K	F act D	D aaa Y aat	K gat F	K tat I	Y ttta E	attgaa
121 2357 136 2353	Q 7 Q 2 F	ĸ	N caa N ttt	S aaa N ata D	A aac T tct S	I aac E gat L	L aca F agt E	K gaa I tta I	E ttc T gaa E	V atc K att	F act D gaa K	D aaa Y aat L	K gat F aaa N	K tat I tta K	Y ttta E aata I	attgaa
121 2357 136 2353 151 2348	Q 7 Q 2 F	ĸ	N caa N ttt	S aaa N ata D	A aac T tct S	I aac E gat L	L aca F agt E aat	K gaa I tta I aac	E ttc T gaa E ata	V atc K att N gta	F act D gaa K	D aaa Y aat L gca	K gat F aaa N att	K tat I tta K gat	Y ttta E aata I aagg	attgaa aaaatt
121 2357 136 2353 151 2348	Q 7 Q 2 F 7	K	N caa N ttt S gac N	S laaa N ata D laga	A aac T tct S aac N	I aac E gat L att	L aca F agt E aat	K gaa I tta I aac V	E ttc T gaa E ata	V atc K att N gta A	F act D gaa K tca	D aaa Y aat L gca	K gat F aaa N att	TAT  tat  ttat  tta  K  gat  V	Y ttta E aata I aaga K	attgaa aaaatt
121 2357 136 2353 151 2348 166 2344	Q 7 Q 2 F 7	K	N caa N ttt S gac N	S laaaa N ata D laga I	A aac T tct S aac N	I aac gat L att N	L aca F agt E aat aat	K gaa I tta I aac V	E ttc gaa E ata tac	V atc K att N gta A	F act D gaa K tca I	D aaa Y aat L gca D ata	K gat F aaa N att	tat I tta K gat V	Y ttta E aata I aagg	attgaa aaaatt gtaaaa
121 2357 136 2353 151 2348 166 2344	Q 7 Q 2 F 7 D 2	K I R	N caa N ttt S gac N aaa	S laaa N lata D laga I ltta G	A aac tct S aac N aaa	I aac gat L att N ggt	L aca F agt aat aat Y	K gaa I tta I aac V agt	E ttc gaa E ata stac	V atc K att N gta A	F act D gaa K tca tca I	D aaa Y aat G D ata	K gat aaa N att att	tat I tta K gat V tct Q	Y ttta E aata I aagg K ttco	attgaa aaaatt gtaaaa
121 2357 136 2353 151 2348 166 2344 181	Q 7 Q 2 F 7 D 2	K I R	N caa N ttt S gac N aaa	S laaa N lata D laga I ltta G	A aac tct S aac N aaa	I aac gat att sggt S ttt	L aca F agt aat aat y cgc	K gaa I tta I aac V agt ata	E ttc gaa E ata s tac aga	V atc A gta gct I aaa	F act D gaa K tca tca I	D aaa Y aat GCa D ata S tta	K gat aaa N att aat F ccc	tat I tta K gat V tct Q	Y ttta E aata I aagg K ttco	attgaa aaaatt gtaaaa cagccg
121 2357 136 2353 151 2348 166 2344 181 2339	Q 7 Q 2 F 7 D 2 K 7	K I R L	N caa N ttt S gac N aaa K att	Saaa Nata Daga Itta Gagt	A aac T S aac N aaa N aag	I aac gat att art ggt ttt	L aca F agt aat aat cgc	K gaa I tta I aac V agt ata K	E ttc gaa E ata s tac aga V	Vatc Katt Ngta Gct Iaaa L	F act D gaa tca tgc tgc gtt	D aaa Y aat GCa D ata S tta	K gat aaa N att aat F ccc	tat I tta K gat tct Caa I	Y ttta E aata I aagg K ttcc P aaaa	attgaa aaaatt gtaaaa cagccg
121 2357 136 2353 151 2348 166 2344 181 2339 196	Q 7 Q 2 F 7 D 2 K 7	K I R L	N caa N ttt S gac N aaa K att	Saaa Nata Daga Itta Gagt	A aac T S aac N aaa N aag	I aac gat att art ggt ttt	aca F agt aat aat cgc Rat	K gaa I taa V aga A ata K tct	E ttc gaa E ata tac agu tcg	Vatc att gta gta gct aaa L	F act D gaa tca tgc tgc gtt	aaa Y aat gca D ata tta Q	K gat aaa N a K t a K t c K t a	tat I tta K gat tct Caa I	Y ttta E aata I aagg K ttcc P aaaa	attgaa aaaatt gtaaaa cagccg
121 2357 136 2353 151 2348 166 2344 181 2339 196 2335	Q 7 Q 2 F 7 D 2 K 7 I 2 N	K I R L	N caa N ttt S gac N aaa K att K aat	Saaa Nata Daga Itta Cagt Cca	A aac tct sac aaa N aag R gta	I aac gat L tt ats tti ats	L aca F agt aat cgc gat gat	K gaa tta taa aaV ata ata tct	E T G E E C G E E C G E E C G E E C G E E C G E E E E	Vatc att gta gta gc aa gat gat	F act gaa tca tg ty gt att	aaa Y aat gca ata ata tta atg	K gat aa N t aa K t C K t t I	tat  ttat  ttat  gav  tcat  cat  cut  cut  cut  cut  cut  cu	Y ttta E aata I aagg K ttco P aaaa A ataa	attgaa aaaatt gtaaaa cagccg

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## Physico-chemical parameters of ORF 77ORF017

1	MTHNIEKRIN	KLKTSGNPKF	KKLDSDIHYL	LKRFEGEKNH	KGFYPKFKQG	EIVFVDFGIN
61	VNKEFSNSHF	AIVMNKNDSN	TEDIVNVIPL	SSKENKKYLK	MNFDLKWEYY	LRLFLNLISA
121	QNNSAILKEV	FDKKYQKNNT	EFITKDYFIE	FISDSLEIEN	KLNKIDRNIN	NIVSAIDKVK
181	KLKGNSYACI	NSFQPISKFR	IRKVLPQKIK	NPVIDSSDIM	LLINRINNNI	LOIPDIR

Number of amino acids:	237
Average molecular weight (Daltons):	27887.38
Mean amino acid weight (Daltons):	117.67
Monoisotopic molecular weight (Daltons):	27869.83
Mean amino acid monoisotopic weight (Daltons):	117.59

## Amino acid composition

Aci d	Symbo	Numb er	%	Average % in Swissprot	Aci d	Symbo l	Numb er	%	Average % in Swissprot
Ala	A	5	2.11%	7.58%	Cys	С	1	0.42%	1.66%
Asp	D	14	5.91%	5.28%	Glu	E	13	5.49%	6.37%
Phe	F	16	6.75%	4.09%	Gly	G	6	2.53%	6.84%
His	Н	4	1.69%	2.24%	Ile	I	29	12.24 %	5.81%
Lys	K	33	13.92 %	5.95%	Leu	L	19	8.02%	9.42%
Met	М	4	1.69%	2.37%	Asn	N	30	12.66 %	4.45%
Pro	P	7	2.95%	4.9%	Gln	Q	6	2.53%	3.97%
Arg	R	8	3.38%	5.16%	Ser	S	17	7.17%	7.12%
Thr	T	5	2.11%	5.67%	Val	V	11	4.64%	6.58%
Тгр	W	1	0.42%	1.23%	Tyr	Y	8	3.38%	3.18%

Number of acidic (negative) amino acids (ED):	27
	11.39%
Number of basic (positive) amino acids (KR):	41
- · · · · · · · · · · · · · · · · · · ·	17.30%
Total charge (KRED):	68
	28.69%
Net charge (KR - ED):	14
	5.91%
Theoritical pI:	10.01
Total linear charge density:	0.30
Average hydrophobicity:	-5.37
Ratio of hydrophilicity to hydrophobicity:	1.41
Percentage of hydrophilic amino acid:	57.81%
Percentage of hydrophobic amino acid:	42.19%
Ratio of %hydrophilic to %hydrophobic:	1.37

WO 00/32825

155

## 77ORF019 sequence

39851		atg												gcagg	aggt
1 M	N	E	Q	I	I	G	S	I	Y	T	L	Α	G	G	
39896		gtt	gtg	ctt	tat	tca	gtt	aaa	gag	att	ttt	agg	tat	tttac	agat
16 V	V	L	Y	S	V	K	E	I	F	R	Y	F	T	D	
39941		tct	aac	tta	caa	cgt	aaa	aaa	atc	aat	tta	gaa	caa	atata	tccg
31 S	N	L	Q	R	K	K	I	N	L	E	Q	I	Y	P	
39986		ata	tat	tta	gat	tgt	ttt	aaa	aag	gct	aaa	aag	atg	attgg	agct
46 I	Y	L	D	С	F	K	K	Α	K	K	M	I	G	Α	
40031		tat	att	att	cca	aca	gaa	.cag	cat	gaa	ttt	tta	gat	tttt	tgat
61 Y	I	I	P	T	E	Q	Н	E	F	L	D	F	F	D	
40076		att	gaa	gtc	ttt	aat	aat	tta	gat	aag	caa	agt	aaa	aaago	gtat
76 I	E	V	F	N	N	L	D	K	Q	S	K	K	Α	Y	
40121		gaa	aat	gtt	att	gga	ttt	aga	caa	atg	att	aat	tta	tcaaa	taga
91 E	N	V	I	G	F	R	Q	M	I	N	L	S	N	R	
40166		gtt	aag	gca	atg	gaa	gat	ttt	aag	atg	agt	ttc	aac	aatga	attt
106 V	K	Α	M	E	D	F	K	M	S	F	N	N	E	F	
40211														atgga	aaca
121 S	T	N	Q	I	F	F	N	P	S	F	V	M	E	T	
40256		att	gct	att	ata	aat	gaa	tat	caa	aaa	gat	ata	tct	tattt	aaaa
136 I	A	I	I	N	E	Y	Q	K	D	I	S	Y	L	K	
40301		aat	ata	att	aat	aaa	atg	aat	gaa	aat	aga	gct	tat	aatca	tatt
151 N	I	I	N	K	M	N	E	N	R	Α	Y	N	H	I	
40346		gat	agt	ttt	atc	act	tca	gag	tac	cga	cga	aaa	ata	aacga	ttat
166 D	S	F	I	T	S	Ε	Y	R	R	K	I	N	D	Y	
40391		aat	ctt	tat	ctt	gat	aaa	ttt	gaa	gaa	cag	ttt	agt	caaaa	gttt
181 N	L	Y	L	D	K	F	E	E	Q	F	S	Q	K	F	
40436		aaa	ata	aac	aga	act	tcg	ata	aaa	gaa	aga	att	att	attaa	ttta
196 K	I	N	R	T	S	I	K	E	R	I	I	I	N	L	
40481		aac	aag	agg	aga	ttt	aaa	tga	40	501					
211 N	K	R	R	F	K	*									

# Physico-chemical parameters of ORF 77ORF019

1	MNEQIIGSIY	TLAGGVVLYS	VKEIFRYFTD	SNLQRKKINL	EQIYPIYLDC	FKKAKKMIGA
61	YIIPTEQHEF	LDFFDIEVFN	NLDKQSKKAY	ENVIGFRQMI	NLSNRVKAME	DFKMSFNNEF
121	STNQIFFNPS	<b>FVMETIAIIN</b>	EYQKDISYLK	NIINKMNENR	AYNHIDSFIT	SEYRRKINDY
181	NLYLDKFEEQ	FSQKFKINRT	SIKERIIINL	NKRRFK		

Number of amino acids:	216
Average molecular weight (Daltons):	26026.06
Mean amino acid weight (Daltons):	120.49
Monoisotopic molecular weight (Daltons):	26009.34
Mean amino acid monoisotopic weight (Daltons):	120.41

## Amino acid composition

Aci d	Symbo l	Numb er	%	Average % in Swissprot	Aci d	Symbo l	Numb er	%	Average % in Swissprot
Ala	Α	7	3.24%	7.58%	Cys	C	1	0.46%	1.66%
Asp	D	10	4.63%	5.28%	Glu	Е	16	7.41%	6.37%
Phe	F	19	8.80%	4.09%	Gly	G	5	2.31%	6.84%
His	Н	2	0.93%	2.24%	Ile	I	28	12.96 %	5.81%
Lys	K	22	10.19 %	5.95%	Leu	L	12	5.56%	9.42%
Met	М	7	3.24%	2.37%	Asn	N	23	10.65 %	4.45%
Pro	P	3	1.39%	4.9%	Gln	Q	10	4.63%	3.97%
Arg	R	11	5.09%	5.16%	Ser	S	13	6.02%	7.12%
Thr	T	7	3.24%	5.67%	Val	V	7	3.24%	6.58%
Ттр	W	0	0.00%	1.23%	Tyr	Y	13	6.02%	3.18%

Number of acidic (negative) amino acids (ED):	26
	12.04%
Number of basic (positive) amino acids (KR):	33
	15.28%
Total charge (KRED):	59
	27.31%
Net charge (KR - ED):	7
	3.24%
Theoritical pI:	9.52
Total linear charge density:	0.28
Average hydrophobicity:	-4.84
Ratio of hydrophilicity to hydrophobicity:	1.37
Percentage of hydrophilic amino acid:	54.17%
Percentage of hydrophobic amino acid:	45.83%
Ratio of %hydrophilic to %hydrophobic:	1.18

## 77ORF043 sequence

2930	4		atg	tat	tac	gaa	ata	ggc	gaa	atc	ata	cgc	aaa	aat	attcate	gtt
1	M	Y	Y	E	I	G	E	I	I	R	K	N	I	H	V	
29349 aacggattcgattttaagctattcattttaaaaggtcatatgggc																
16	N	G	F	D	F	K	L	F	I	$\mathbf{L}$	K	G	H	M	G	
2939	4		ata	tca	ata	caa	gtt	aaa	gat	atg	aac	aac	gta	cca	attaaa	at
31	I	S	I	Q	V	K	D	М	N	N	V	P	I	K	H	
29439 gcttatgtcgtagatgagaatgacttagatatggcatcagactta																
2773	,		900	Luc	900	5	3	2-2	~~~			J - · · · ·	5			
46	A	Y	V	. V	D	E	N	D	L	D	M	A	s	D	L	. <b>.</b> .
	A	Y	V	. <b>V</b>	D	E	N	D	L	D	M	Α	s	D	L acagac	
46	A	Y	V	. <b>V</b>	D	E	N	D	L	D	M	Α	s	D	L	
46 2948	A 4 F	N	V ttt Q	V aac A	D caa I	E gca D	N ata E	D gate W	L gaa I	D tgg: E	M att E	A gaaq N	S gag T	D aaca D	L acagac	

## Physico-chemical parameters of ORF 77ORF043

MYYEIGEIIR KNIHVNGFDF KLFILKGHMG ISIQVKDMNN VPIKHAYVVD ENDLDMASDLFNQAIDEWIE ENTDEQDRLI NLVMKW

61

Number of amino acids:	86
Average molecular weight (Daltons):	10186.68
Mean amino acid weight (Daltons):	118.45
Monoisotopic molecular weight (Daltons):	10180.02
Mean amino acid monoisotopic weight (Daltons):	118.37

## Amino acid composition

Aci d	Symbo	Numb er	%	Average % in Swissprot	Aci d	Symbo l	Numb er	%	Average % in Swissprot
Ala	Α	3	3.49%	7.58%	Cys	С	0	0.00%	1.66%
Asp	D	9	10.47 %	5.28%	Glu	E	7	8.14%	6.37%
Phe	F	4	4.65%	4.09%	Gly	G	4	4.65%	6.84%
His	н	3	3.49%	2.24%	Ile	I	11	12.79 %	5.81%
Lys	K	6	6.98%	5.95%	Leu	L	6	6.98%	9.42%
Met	M	5	5.81%	2.37%	Asn	N	8	9.30%	4.45%
Pro	P	1	1.16%	4.9%	Gln	Q	3	3.49%	3.97%
Arg	R	2	2.33%	5.16%	Ser	S	2	2.33%	7.12%
Thr	T	1	1.16%	5.67%	Val	V	6	6.98%	6.58%
Trp	W	2	2.33%	1.23%	Туг	Y	3	3.49%	3.18%

Number of acidic (negative) amino acids (ED):	16
Number of basic (positive) amino acids (KR):	18.60% 8
	9.30%
Total charge (KRED):	24
·	27.91%
Net charge (KR - ED):	-8
9.30%	
Theoritical pI:	4.38
Total linear charge density:	0.30
Average hydrophobicity:	-2.80
Ratio of hydrophilicity to hydrophobicity:	1.19
Percentage of hydrophilic amino acid:	48.84%
Percentage of hydrophobic amino acid:	51.16%
Ratio of %hydrophilic to %hydrophobic:	0.95

159

#### 77ORF102 sequence

- 12.4

# Physico-chemical parameters of ORF 77ORF102

1 MSNIYKSYLV AVLCFTVLAI VLMPFLYFTT AWSIAGFASI ATFMYYKECF FKE

Number of amino acids:	53
Average molecular weight (Daltons):	6155.42
Mean amino acid weight (Daltons):	116.14
Monoisotopic molecular weight (Daltons):	6151.07
Mean amino acid monoisotopic weight (Daltons):	116.06

### Amino acid composition

Aci d	Symbo	Numb er	%	Average % in Swissprot	Aci d	Symbo I	Numb er	%	Average % in Swissprot
Ala	A	6	11.32 %	7.58%	Cys	С	2	3.77 %	1.66%
Asp	D	0	0.00%	5.28%	Glu	E	2	3.77 %	6.37%
Phe	F	7	13.21 %	4.09%	Gly	G	1	1.89 %	6.84%
His	Н	0	0.00%	2.24%	Ile	I	4	7.55 %	5.81%
Lys	K	3	5.66%	5.95%	Leu	L	5	9.43 %	9.42%
Met	М	3	5.66%	2.37%	Asn	N	1	1.89 %	4.45%
Pro	P	1	1.89%	4.9%	Gln	Q	0	0.00 %	3.97%
Arg	R	0	0.00%	5.16%	Ser	s	4	7.55 %	7.12%
Thr	Т	4	7.55%	5.67%	Val	V	4	7.55 %	6.58%
Trp	w	1	1.89%	1.23%	Tyr	Y	5	9.43 %	3.18%

(negative) amino acids (ED):	
3.77%	
3	
5.66%	
5	
9.43%	
1	
1.89%	
8.18	
0.13	
10.81	
0.40	
28.30%	
71.70%	

WO 00/32825

161

Ratio of %hydrophilic to %hydrophobic:

0.39

162

#### 77ORF104 sequence

atggtaaccaaagaatttttaaaaactaaacttgagtgttcagat

M V T K E F L K T K L E C S D

atgtacgctcagaaactcatagatgaggcacagggcgatgaaaat

M Y A Q K L I D E A Q G D E N

aggttgtacgacctatttatccaaaaacttgcagaacgtcataca

R L Y D L F I Q K L A E R H T

34528 cgcccgctatcgtcgaatattaa 34551

46 R P A I V E Y \*

- \_\_\_\_\_

# Physico-chemical parameters of ORF 77ORF104

1 MVTKEFLKTK LECSDMYAQK LIDEAQGDEN RLYDLFIQKL AERHTRPAIV EY

Number of amino acids:	52
Average molecular weight (Daltons):	6193.13
Mean amino acid weight (Daltons):	119.10
Monoisotopic molecular weight (Daltons):	6189.12
Mean amino acid monoisotopic weight (Daltons):	119.02

## Amino acid composition

Aci d	Symbo l	Numb er	%	Average % in Swissprot	Aci d	Symbo l	Numb er	%	Average % in Swissprot
Ala	A	4	7.69 %	7.58%	Cys	С	1	1.92%	1.66%
Asp	D	4	7.69 %	5.28%	Glu	Е	6	11.54 %	6.37%
Phe	F	2	3.85 %	4.09%	Gly	G	1	1.92%	6.84%
His	Н	1	1.92 %	2.24%	Ile	I	3	5.77%	5.81%
Lys	K	5	9.62 %	5.95%	Leu	L	6	11.54 %	9.42%
Met	М	2	3.85 %	2.37%	Asn	N	1	1.92%	4.45%
Pro	P	1	1.92	4.9%	Gln	Q	3	5.77%	3.97%
Arg	R	3	5.77 %	5.16%	Ser	S	1	1.92%	7.12%
Thr	Т	3	5.77 %	5.67%	Val	V	2	3.85%	6.58%
Trp	w	0	0.00 %	1.23%	Tyr	Y	3	5.77%	3.18%

Number of acidic (negative) amino acids (ED):	10
	19.23%
Number of basic (positive) amino acids (KR):	8
-	15.38%
Total charge (KRED):	18
	34.62%
Net charge (KR - ED):	-2 -
3.85%	
Theoritical pI:	5.03
Total linear charge density:	0.38
Average hydrophobicity:	-5.81
Ratio of hydrophilicity to hydrophobicity:	1.47
Percentage of hydrophilic amino acid:	53.85%
Percentage of hydrophobic amino acid:	46.15%

164

Ratio of %hydrophilic to %hydrophobic:

1.17

165

## 77ORF182 sequence

2926	29268 atgttcaatataaaacgaaaacggaggaagtcaagatgtattac															
1	M	F	N	I	K	R	K	T	Ε	Ε	V	K	M	Y	Y	
29313 gaaataggcgaaatcatacgcaaaaatattcatgttaacggattc												ggattc				
16	Ε	I	G	E	I	I	R	K	N	I	H	V	N	G	F	
29358 gattttaagctattcattttaaaaggtcatatgggcatatcaata																
31	D	F	K	L	F	I	L	K	G	H	M	G	I	S	I	
2940	)3		caa	gtt	aaa	gat	atg	aac	aac	gta	cca	att	aaa	cat	gcti	tatgtc
46	Q	V	K	D	M	N	N	V	P	I	K	H	Α	Y	V	
2944	18		gta	gat	gag	aat	gac	tta				tca	gac	tta	ttta	aaccaa
61	V	D	E	N	D	L	D	M	A	S	D	L	F	N	Q	
2949	93		gca	ata	gat	gaa	tgg	att	gaa	gag	aac	aca	gac	gaa	cag	gacaga
76	A	I	D	E	W	I	Ε	E	N	${f T}$	D	E	Q	D	R	
2953	38		cta	att	aac	tta	gtc	atg	aaa	tgg	tag	29	564			
91	L	I	N	L	V	M	K	W	*							

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## Physico-chemical parameters of ORF 77ORF182

1 MFNIKRKTEE VKMYYEIGEI IRKNIHVNGF DFKLFILKGH MGISIQVKDM NNVPIKHAYV

VDENDLDMAS DLFNQAIDEW IEENTDEQDR LINLVMKW

Number of amino acids: 98

Average molecular weight (Daltons): 11691.50

Mean amino acid weight (Daltons): 119.30

Monoisotopic molecular weight (Daltons): 11683.84

Mean amino acid monoisotopic weight (Daltons): 119.22

#### Amino acid composition

Aci d	Symbo	Numb er	%	Average % in Swissprot	Aci d	Symbo l	Numb er	%	Average % in Swissprot
Ala	A	3	3.06 %	7.58%	Cys	С	0	0.00%	1.66%
Asp	D	9	9.18 %	5.28%	Glu	E	9	9.18%	6.37%
Phe	F	5	5.10 %	4.09%	Gly	G	4	4.08%	6.84%
His	Н	3	3.06 %	2.24%	Ile	I	12	12.24 %	5.81%
Lys	К	9	9.18 %	5.95%	Leu	L	6	6.12%	9.42%
Met	М	6	6.12 %	2.37%	Asn	N	9	9.18%	4.45%
Pro	P	1	1.02 %	4.9%	Gln	Q	3	3.06%	3.97%
Arg	R	3	3.06 %	5.16%	Ser	s	2	2.04%	7.12%
Thr	Т	2	2.04 %	5.67%	Val	V	7	7.14%	6.58%
Trp	w	2	2.04 %	1.23%	Tyr	Y	3	3.06%	3.18%

Number of acidic (negative) amino acids (ED):	18
	18.37%
Number of basic (positive) amino acids (KR):	12
	12.24%
Total charge (KRED):	30
	30.61%
Net charge (KR - ED):	-6 -
6.12%	
Theoritical pI:	4.76 ~
Total linear charge density:	0.33
Average hydrophobicity:	-3.89
Ratio of hydrophilicity to hydrophobicity:	1.28

167

Percentage of hydrophilic amino acid:	51.02%
Percentage of hydrophobic amino acid:	48.98%
Ratio of %hydrophilic to %hydrophobic:	1.04

168

#### Table 5

BLASTP 2.0.8 [Jan-05-1999] Query= sid|100017|lan|770RF017 Phage 77 ORF |23269-23982|-3 (237 letters)

Database: nr

393,678 sequences; 120,452,765 total letters

Sequences producing significant alignments:	Score (bits)	E Value
	,	
gi 4493986 emb CAB39045.1  (AL034559) predicted using hexExon;	41	0.010
gi 730607 sp P23250 RPI1_YEAST NEGATIVE RAS PROTEIN REGULATOR P	38	0.053
gi 3097044 emb CAA75299  (Y15035) K1R [Cowpox virus]	38	0.090
gi 2146245 pir  S73794 hypothetical protein H91_orf180 - Mycopl	38	0.090
gi 83910 pir  S04682 ribosomal protein varl - yeast (Candida gl	37	0.15
gi   133135   sp   P21358   RMAR_CANGA MITOCHONDRIAL RIBOSOMAL PROTEIN	37	0.15
gi 2128843 pir  H64475 hypothetical protein MJ1409 - Methanococ	36	0.20
gi 5107017 gb AAD39926.1 AF126285_2 (AF126285) RNA polymerase (	36	0.35
gi 2146210 pir  S73342 hypothetical protein E07_orf166 - Mycopl	35	0.60

Database: swissprot 79,449 sequences; 28,874,452 total letters

Sequ	iences l	producing si	gnificant alignments:	Score (bits)	E Value
вp	P23250	RPI1_YEAST	NEGATIVE RAS PROTEIN REGULATOR PROTEIN.	38	0.014
sp	P21358	RMAR_CANGA	MITOCHONDRIAL RIBOSOMAL PROTEIN VAR1.	37	0.040
sp	Q21444	LDLC_CAEEL	LDLC PROTEIN HOMOLOG.	34	0.35
sp	P27240	RFAY_ECOLI	LIPOPOLYSACCHARIDE CORE BIOSYNTHESIS PROT.	. 33	0.46
sp	P53192	YGC0_YEAST	HYPOTHETICAL 27.1 KD PROTEIN IN ALK1-CKB1.	. 33	0.60
sp	P32908	SMC1_YEAST	CHROMOSOME SEGREGATION PROTEIN SMC1 (DA-B.	. 33	0.60
sp	P54683	TAGE DICDI	PRESTALK-SPECIFIC PROTEIN TAGB PRECURSOR .	32	0.78
sp	Q03100	CYAA_DICDI	ADENYLATE CYCLASE, AGGREGATION SPECIFIC (.	32	0.78

169

BLASTP 2.0.8 [Jan-05-1999]

Query= sid|100019|lan|770RF019 Phage 77 ORF|39851-40501|2 (216 letters)

Database: nr

373,355 sequences; 114,214,446 total letters

·	Score	E Value
gi 3341966 dbj BAA31932  (AB009866) orf 59 [bacteriophage phi PVL]	437	e-122
gi 2689911 (AE000792) B. burgdorferi predicted coding region BB	38	0.058
gi 1171589 emb CAA64574  (X95275) frameshift [Plasmodium falcip	37	0.10
qi 4493986 emb CAB39045.1  (AL034559) predicted using hexExon;	36	0.23
gi 141257 sp P18019 YPI9 CLOPE HYPOTHETICAL 14.5 KD PROTEIN (OR	36	0.29
gi 133412 sp P27059 RPOB ASTLO DNA-DIRECTED RNA POLYMERASE BETA	35	0.51
gi 3122231 sp Q58851 HISX METJA HISTIDINOL DEHYDROGENASE (HDH)	35	0.51
gi 3649757 emb CAB11106.1 (Z98547) predicted using hexExon; MA	34	0.66
gi 2688313 (AE001146) sensory transduction histidine kinase, pu	34	0.87

Database: swissprot

79,449 sequences; 28,874,452 total letters

Sequences	producing a	significant alignments:	Score (bits)	E Value
sp P18019	YPI9_CLOPE	HYPOTHETICAL 14.5 KD PROTEIN (ORF9).	36	0.079
sp Q58851	HISX_METJA	HISTIDINOL DEHYDROGENASE (EC 1.1.1.23) (H.	35	0.14
sp   P27059	RPOB_ASTLO	DNA-DIRECTED RNA POLYMERASE BETA CHAIN (E.	35	0.14
sp Q02224	CENE_HUMAN	CENTROMERIC PROTEIN E (CENP-E PROTEIN).	34	0.31
sp P04931	ARP_PLAFA	ASPARAGINE-RICH PROTEIN (AG319) (ARP) (FRA	. 33	0.53
sp P18011	IPAB_SHIFL	62 KD MEMBRANE ANTIGEN.	32	0.69
sp P18709	VTA2_XENLA	VITELLOGENIN A2 PRECURSOR (VTG A2) [CONTA	. 32	0.90
sp Q64409	CP3H_CAVPO	CYTOCHROME P450 3A17 (EC 1.14.14.1) (CYPI	. 32	0.90
sp P21358	RMAR_CANGA	MITOCHONDRIAL RIBOSOMAL PROTEIN VAR1.	32	0.90
sp Q03945	IPAB_SHIDY	62 KD MEMBRANE ANTIGEN.	32	1.2

- ...--

170

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BLASTP 2.0.8 [Jan-05-1999]
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Query= sid|100043|lan|770RF043 Phage 77 ORF|29304-29564|3 (86 letters)

Database: nr

373,355 sequences; 114,214,446 total letters

Sequences producing significant alignments:	Score	E Value
gi 3341947 dbj BAA31913  (AB009866) orf 39 (bacteriophage phi PVI gi 744518 prf  2014422A FKBP-rapamycin-associated protein [Homo. gi 1169736 sp P42346 FRAP_RAT FKBP-RAPAMYCIN ASSOCIATED PROTEIN. gi 1169735 sp P42345 FRAP_HUMAN FKBP-RAPAMYCIN ASSOCIATED PROTE. gi 3282239 (U88966) rapamycin associated protein FRAP2 [Homo sa. gi 3875402 emb CAA98122  (Z73906) cDNA EST EMBL:D64544 comes fr. gi 1084792 pir  S54091 hypothetical protein YPR070w - yeast (Sa.	. 32 . 32 . 32 . 32 . 31	0.84 0.84
Database: swissprot		

base: swissprot
 79,449 sequences; 28,874,452 total letters

Sequences	producing significant alignments:	Score (bits)	E Value
sp P42345	FRAP HUMAN FKBP-RAPAMYCIN ASSOCIATED PROTEIN (FRAP)	32	0.24
sp P42346	FRAP RAT FKBP-RAPAMYCIN ASSOCIATED PROTEIN (FRAP) (R.	. 32	0.24
sp P34554	YNP1 CAEEL HYPOTHETICAL 42.2 KD PROTEIN T05G5.1 IN C.	. 28	3.5
sp Q24118	LIO_DROME LINOTTE PROTEIN.	28	3.5
sp P80034	ACH2_BOMMO ANTICHYMOTRYPSIN II (ACHY-II).	28	3.5
sp   P22922	ALAT BOMMO ANTITRYPSIN PRECURSOR (AT).	28	3.5
sp Q44363	TRAA AGRT6 CONJUGAL TRANSFER PROTEIN TRAA.	28	3.5
sp P38255	YBU5_YEAST HYPOTHETICAL 51.3 KD PROTEIN IN PHO5-VPS1.	27	6.0
sp P55822	SH3B HUMAN SH3BGR PROTEIN (21-GLUTAMIC ACID-RICH PRO.	27	7.9
sp   Q58482	YA82_METJA HYPOTHETICAL PROTEIN MJ1082.	27	7.9
sp   P34252	YKK8 YEAST HYPOTHETICAL 52.3 KD PROTEIN IN HAP4-AAT1.	27	7.9

171

BLASTP 2.0.8 [Jan-05-1999]

Query= sid|100102|lan|770RF102 Phage 77 ORF|29051-29212|2 (53 letters)

Database: nr

373,355 sequences; 114,214,446 total letters

Sequences producing significant alignments: (bit		E Value
gi 3341946 dbj BAA31912  (AB009866) orf 38 [bacteriophage phi PVL] gi 4325288 gb AAD17315  (AF123593) voltage-dependent sodium cha gi 2649684 (AE001040) A. fulgidus predicted coding region AF092	28	3e-20 7.1 9.3
Database: swissprot 79,449 sequences; 28,874,452 total letters		
Sco Sequences producing significant alignments: (bit		E Value
	26	7.1
sp P04775 CIN2 RAT SODIUM CHANNEL PROTEIN, BRAIN II ALPHA SUBU sp P42619 YQJF_ECOLI HYPOTHETICAL 17.2 KD PROTEIN IN EXUR-TDCC	26	9.2

172

BLASTP 2.0.8 [Jan-05-1999]

Query= sid|100104|lan|770RF104 Phage 77 ORF|34393-34551|1 (52 letters)

Database: nr

373,355 sequences; 114,214,446 total letters

Sequences producing significant alignments:	Scor (bits	E E Value
gi 2315523 (AF016452) similar to the leucine-rich domains found gi 4377168 gb AAD18990  (AE001666) CT711 hypothetical protein [gi 3882171 dbj BAA34445  (AB018268) KIAA0725 protein [Homo sapi	2	5.4

Database: swissprot

79,449 sequences; 28,874,452 total letters

Sequences producing s	ignificant alignments:	Score (bits)	E Value
sp P04879 RRPP_VSVIG	RNA POLYMERASE ALPHA SUBUNIT (EC 2.7.7.48.	27	5.4
sp P04880 RRPP VSVIM	RNA POLYMERASE ALPHA SUBUNIT (EC 2.7.7.48.	27	5.4
sp Q13946 CN7A HUMAN	HIGH-AFFINITY CAMP-SPECIFIC 3',5'-CYCLIC .	26	7.1
sp P35381 ATPA DROME	ATP SYNTHASE ALPHA CHAIN, MITOCHONDRIAL P.	26	9.3
sp P54659 MVPB_DICDI	MAJOR VAULT PROTEIN BETA (MVP-BETA).	26	9.3
sp P40397 YHXC BACSU	HYPOTHETICAL OXIDOREDUCTASE IN APRE-COMK .	26	9.3

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173

BLASTP 2.0.8 [Jan-05-1999]

Query= sid|122748|lan|770RF182 Phage 77 ORF|29268-29564|3 (98 letters)

Database pr

Database: nr 393,678 sequences; 120,452,765 total letters		
Sequences producing significant alignments:	Score (bits)	_
gi 3341947 dbj BAA31913.1  (AB009866) orf 39 (bacteriophage phigilo84792 pir  S54091 hypothetical protein YPR070w - yeast (Sagi 1169736 sp P42346 FRAP_RAT FKBP-RAPAMYCIN ASSOCIATED PROTEIN. gi 744518 prf  2014422A FKBP-rapamycin-associated protein [Homo.gi 5051381 emb CAB44736.1  (AL049653) dJ647M16.2 (FK506 binding.gi 4826730 ref NP_004949.1 pFRAP1  FK506 binding protein 12-rap.gi 3282239 (U88966) rapamycin associated protein FRAP2 [Homo sa.	. 35 . 32 . 32 . 32	8e-46 0.13 1.1 1.1 1.1 1.1
Database: swissprot 79,909 sequences; 29,054,478 total letters		
Sequences producing significant alignments:	Score (bits)	
sp P42345 FRAP_HUMAN FKBP-RAPAMYCIN ASSOCIATED PROTEIN (FRAP) . sp P42346 FRAP_RAT FKBP-RAPAMYCIN ASSOCIATED PROTEIN (FRAP) (R. sp P40557 YIA5_YEAST PUTATIVE DISULFIDE ISOMERASE YIL005W PREC. sp Q24118 LIO_DROME LINOTTE PROTEIN. sp P80034 ACH2_BOMMO SP P80534 ACH2_BOMMO ANTICHYMOTRYPSIN II (ACHY-II). sp P34554 YNP1_CAEEL HYPOTHETICAL 42.2 KD PROTEIN TO5G5.1 IN C. sp P22922 A1AT BOMMO ANTITRYPSIN PRECURSOR (AT).	32 29 28 28 28	0.29 0.29 3.3 4.4 4.4 4.4

Table 6

1st position (5' end)	2nd position				3rd position (3' end)
U	Phe	Ser	Tyr	Cys	U
	Phe	Ser	Туг	Cys	l c
	Leu	Ser	Stop	Stop	A
	Leu	Ser	Stop	Trp	
С	Leu	Pro	His	Arg	U
	Leu	Pro	His	Arg	c
	Leu	Pro	Gln	Arg	Α
	Leu	Pro	Gln	Arg	G
Α	lle	Thr	Asn	Ser	U
	lle	Thr	Asn	Ser	С
	lle	Thr	Lys	Arg	A
	Met_	Thr	Lys	Arg	G
G	Val	Ala	Asp	Gly	U
	Val	Ala	Asp	Gly	С
	Val	Ala	Glu	Gly	A
	Val	Ala	Glu	Glv	G

175

Table 7

Bacteriophage 3A, complete genome sequence

```
caaacgctag caacgcggat aaatttttca tgaaaggggg tctttatatg aagttaacaa aaaaacagct
        aaaagaatat atagaagatt acaaaaaatc tgatgacata ttaattaatt tgtatataga aacatatgaa
71
        ttttattgtc ggttaagaga tgaacttaaa aatagtgatt taatgataga gcatacaaac aaggctggtg
141
        cqaqcaatat tattaagaat ccattaagca tagaactgac aaaaacagtt caaacactaa ataacttact
211
        caagtctatg ggtttaactg cagcacaaag aaaaaagata gttcaagaag aaggtggatt cggtgactat
281
        taaagtttta aatgaacctt caccaaaact attaacaaca tggtatgcag agcaagtcac tcaagggaaa
351
421
        ataaaaacaa gcaaatatgt tagaaaagaa tgtgagagac atcttagata tctagaaaat ggaggtaaat
491
       gggtatttga tgaagaatta gcgcatcgtc ctattcgatt tatagaaaag ttttgtaaac cttccaaagg
561
        atctaaacgt caacttgtat tacagccatg gcaacatttt attatcggca gtttgtttgg ttgggttcat
631
        aaagaaacaa aactgcgcag gtttaaagaa gctttgatat ttatggggcg aaaaaatggt aaaacaacca
701
        ctatttctgg ggttgctaac tatgctgtat Cacaagatgg agaaaatggt gcagaaattc atttgttagc
771
        anacqtaatg aaacaagcta ggattctatt tgatgaatct aaggcgatga ttaaagctag cccaaagctt
841
       gataaaaatt tcagaacatt aagagatgaa atccattatg acgcaacgat atcaaaaatt atgccccaag
911
        catcagatag cgataagtta gatggattga atacacacat ggggattttt gatgaaattc atgaatttaa
981
        agactataaa ttgatttcag ttataaaaaa ctcaagagct gcaaggttac aacctcttct catctacatt
1051
       acgacagcag ggtatcaatt agatggtcca cttgttgata tggtagaagc gggaagagac accttagatc
1121
       aaatcataga agacgaaaga actttttatt atttagcatc tttggatgat gacgatgata ttaatgattc
1191
       gtcgaactgg ataaaagcaa atcccaactt aggtgtctct ataaatttag atgagatgaa agaagagtgg
1261
       gaaaaagcta agagaacacc agctgaacgt ggagatttta taaccaaaag gtttaatatc tttgctaata
1331
        atgacgagat gagttttatt gattacccaa cactccaaaa aaataatgaa attgtttctt tagaagagct
1401
       ggaaggcaga ccgtgcacga ttggttatga tttatcagaa acagaggact ttacagccgc gtgtgctact
1471
        tttgcgttag ataatggtaa agttgcagtt ttatcgcatt catggattcc taagcacaaa gttgaatatt
1541
       ctaacgaaaa aataccctat agagaatggg aagaagatgg cttattaaca gtgcaagata agccttatat
1611
       tgactaccaa gatgttttaa attggataat taagatgaat gagcattatg tagtagaaaa aattacttat
1681
       gatagagcga acgcattcaa actaaatcaa gagttaaaaa attacgggtt tgaaacggaa gaaacaagac
1751
        aaggagettt gaeettgage eetgeattga aggatttaaa agaaatgttt ttagatggga aaataatatt
1821
       taataataat cctttaatga aatggtatat caataatgtt cagttgaaac tagacagaaa cggaaactgg
1891
       ttgccgtcta agcaaagcag atatcgtaaa atagatggct ttgcagcatt tttaaacaca tatacagata
1961
       ttatgaataa agttgttttt gatagtggtg aaggaaacat agagtttatt agtattaaag acataatgcg
2031
       ttaaggaggt gaatgttatc gcaaaagaga atattgtcac acgcataaag aaaaaattga tagacaattg
2101
       gattgatcag tcaacttcta agctttatga ctttagccca tggaaaaata gatctttttg gggtgtaatt
2171
       aataatacgc ttgaaactaa tgaaacgata ttttcagcta ttacaaagtt atctaattcg atggctagtt
2241
       tgcccttgaa aatgtatgaa gattataaag tagttaatac agaagtatet gatttactta cagtgtcacc
2311
       gaataattot otgagoagtt ttgattttat taatoaaatt gaaacaatca gaaatgaaaa aggtaatgoa
2381
       tatgtgctaa ttgaacgaga catctatcat caaccatcaa agcttttctt attaaatcca gatgttgttg
2451
       aaatgttaat tgaaaaccaa tcacgtgaac tttattattc cattcatgct gcaactggaa ataaattgat
2521
       tgttcataat atggacatgt tgcattttaa acacatcgtg gcatctaata tggtgcaagg cattagtccg
2591
       attgatgtgt tgaagaatac aactgatttt gataatgcag taagaacctt taatcttaca gaaatgcaaa
2661
       aacctgattc tttcatgctt aaatatggtt ccaatgtagg taaagaaaaa aggcagcaag tgttagaaga
2731
       tttcaaacag tactatgaag aaaacggtgg aatattattc caagagcctg gtgttgaaat cgaaccgtta
2801
       cctaaaaaat atgtctctga agatatagtg gcaagcgaga atttaacaag agaaagagta gctaacgttt
2871
       ttcaattgcc ctcagtattc ttaaatgcaa gatcaaatac aaatttcgcg aaaaatgaag agttaaacag
2941
       attttacttg cagcatacct tattgccaat cgtcaaacag tatgaagaag aatttaatcg gaaactactt
3011
       actaaaacag acagagaaaa aaataggtat tttaaattta acgttaaatc ttatttaagg gctgatagtg
3081
       caacacaagc agaagtgtac tttaaagcag ttcgtagtgg ttactacact ataaatgaca ttagagagtg
3151
       ggaagattta ccaccagttg aaggtggaga taagccgcta ataagcggtg atttataccc aattgacacg
3221
       ccacttgaat taagaaaatc tttgaaaggt ggtgataaaa atgtcaatga aagctaagta ttttcaaatg
3291
       aaaagaaaat caaaaagtaa aggtgaaata tttatttatg gtgatattgt aagtgataaa tggtttgaaa
3361
       gtgatgtaac tgctacagat ttcaaaaata aactagatga actaggagac atcagtgaaa tagatgttca
3431
       tataaattca totggaggca gtgtatttga agggcatgca atatacaata tgctaaaaat gcatcotgca
3501
       aaaattaata totatgtoga tgoottagog goatcaattg otagtgttat ogotatgagt ggtgacacta
3571
       tttttatgca caaaaatagt tttttaatga ttcataattc atgggttatg actgtaggta atgcagaaga
3641
       gttaagaaag acagcggatt tacttgaaaa aacagatgct gttagtaatt cagcttattt agataaagca
3711
       aaagatttag atcaagaaca cttaaaacag atgttagatg cagaaacttg gcttactgca gaagaagcct
       tgtctttcgg cttgatagat gaaattttag gagctaatga aataactgct agtatctcta aagagcaata taagcgtttc gagaacgtcc cagaagattt aaagaaagat gtagacaaaa tcactaaaat cgatgatgta
3781
3851
3921
       gatacgtttg aattggttga aacacctaaa gaaagtatgt cactagaaga aaaagaaaaa agagaaaaaa
3991
       ttaaaacgcga atgcgaaatt ttaaaaaatga caatgagtta ttaggaggaa atgaaatgcc gacattatat
4061
       gaattaaaac aatccttagg tatgattgga caacaattaa aaaataaaaa tgatgaattg agtcagaaag
4131
       caacagaccc aaatattgat atggaagaca tcaaacaact agaaacagaa aaagcaggct tacaacaaag
4201
       atttaacatt gttgaaagac aagtaaaaga cattgaagaa aaagaaaaag cgaaagttaa agacacagga
       gaagettate aatetttaaa tgateatgag aagatggtta aagetaagge agagtttat egteacgega
4271
4341
       ttttaccaaa tgaatttgaa aaaccttcaa tggaggcaca acgtttatta cacgctttac caacaggtaa
4411
       tgattcaggt ggtgataagc tcttaccaaa aacactttct aaagaaattg tttcagaacc atttgctaaa
4481
       aaccaattac gtgaaaaagc tcgtctaact aacattaaag gtttagagat tccaagagtt tcatatactt
4551
       tagacgatga tgacttcatt acagatgtag aaacagcaaa agaattaaaa ttaaaaggtg atacagttaa
4621
       atteactact aataaattea aagtatttge tgeaatttea gatactgtaa tteatggate agatgtagat
4691
       ttagtaaact gggttgaaaa cgcactacaa tcaggtctag cagctaaaga acgtaaagat gccttagcag
```

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aatgttcaag aaatggagtg aagcataatg agcgtaatca gtaacagaaa agtagatatg aacgaagcgc 34651 aagacaatgt taagcaacca gcgcactaca catacggcga cattgaaatt atagatttta tcgaacaggt 34721 34791 tacggcacag tatccacctc aactagcatt cgcaataggt aatgcaataa aatacttgtc tagagcacct 34861 ttaaagaatg gtcatgagga tttagcaaag gcgaagtttt acgtccaaag agcttttgac ttgtgggagt 34931 gatgaccatg acagatageg catgtaaaga atacttaaac caatttttcg gatctaagag atatctgtat 35001 caggataacg aacgagtggc acatatccat gtagtgaatg gcacttatta ctttcacggg catatcgtac 35071 caggctggca aggcgtgaaa aagacatttg atacagcgga agagctcgaa acatatataa agcaacatgg tttggaatac gaggaacaga agcaactaac tttattttaa ggagatagaa atgatgaaaa tcaaaagttga 35141 35211 aaaaataatg aaaatagacg aattaattaa gtgggcgcga gaaaatccgg agctatcatt tggcagaaaa 35281 tattatacaa cagacaaaaa tgatgaaaac tttatttact tcggtgtttt taaaaattgt tttaaaataa 35351 gcgattttat attagttaat gctactttta gtgtcaaagt tgaagaagaa gtaaccgaag aaactaagtt 35421 tgataggttg tttgaagtgt acgagattca agaaggagtc tataaatctg catcatatga gaatgctagt 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WO 00/32825 PCT/IB99/02040

182

ggagcaaaag gcttcggaag tagcggagtg taaagacata ttagatcgag tcaaggaggt tttggggaag 36681 tgagtgacat gttagaaata tttttcatag ggtttggtgt ttatctattt tgtcgcatag gtattatttt 36751 totcaagagt aaaaagacta tacacacaaa cotatatgaa atgttgttga ttgctactat ctttgtgaca 36821 tctacatttg ctgataaaca tcaaaagacg catatcttaa tagcattttt agtaatgttt tttatgagta 36891 ageteaaaca agtteaaggg agetatgagg aatgacacaa tacetagtea caacatttaa agatteaaca 36961 ggacgtaagc atacacacat aactaaagct aagagcaatc aaaggtttac agttgttgat gcggagagta 37031 aagaagaagc gaaagagaag tacgaggcac aagttaaaag aaatgcagtt attaaattag ggcagttgtt 37101 tgaaaatata agggagtgtg ggaaatgact aaacaaatac taagattatt attcttacta gcgatgtatg 37171 agctaggcaa gtatgtaact gagcaagtat atattatgat gacggctaat gatgatgcag aggcgccgag 37241 tgactttgaa aaaatcagag ctgaagtttc atggtaatag ctattatcat ttttgaatta attatattaa 37311 tgtgtttagc aatagcactg gaggtgttgt aaatatgtgg attgtcattt caatgtttt atctatattt ttattgatct tgttaagtag catttctcat aagatgaaaa ccatagaagc attggagtat atgaatgctt 37381 37451 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tgcaagatta tatagacaag gacaaaatca tacgactatt attcatcaca tcatgaccga taacacaata

WO 00/32825 PCT/IB99/02040

183

42001	gatcaaagag	tatataaagc	tttacaaaat	aaagaactaa	cgcaagaaga	attgatgaaa	gctattaaag
42071	caagaatagc	taagcataag	taatggaggt	ataagatggg	aaaggcgtca	tatgatatta	agccaggaac
42141	atttaaatat	attgaatcag	aaatatataa	tttaaatgag	aacaagaaag	agataaatag	attgagaatg
42211	gagatactta	acccaacgaa	agaactagac	accaacattg	tgtatggacc	gttacaaaaa	ggagagccag
42281	ttagaacaac	tgagttaatg	gcgacaaggt	tattgactaa	taagatgtta	cgtaacttag	aagagatggt
42351	tgaagcagtt	gaaagtgagt	acttaaagtt	acctgaagat	cataagaaag	taataaggtt	aaagtattgg
42421						gcatcgcaat	
42491	caatacgaaa	gaactttgtt	aaagcgatag	cgtatcatgc	aggtatcaaa	taacattgtg	caaagattgt
42561						aaagttatct	
42631						aatcaaagag	
42701						gattggttct	
42771						gtcaaatgtg	
42841						agattttaac	
42911						gcaaatgata	
42981						aataaaattt	
43051			tttttcgccg				-

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Table 8

# Bacteriophage 3A ORFs list

SID	LAN	PRA	POS	a.a.	RBS sequence	STA	STO
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100380	3AORF002	2	3766740114	815	tttaaaataatgaaaggagccgaac	atg	taa
100381	3AORF003	1	3218834149	653	ttaaagaaattgaggtgtcaagaat	ttg	tag
100382	3AORF004	3	1745719370	637	gctattttattagaaaggaaggtgc	att	taa
100383	3AORF005	1	3342034	566	agaaaaaagatagttcaagaagaag	gtg	taa
100384	3AORF006	1	1557117154	527	cttttatttataggtaggtgattta	atg	taa
100385	3AORF007	2	1933720836	499	atgatagtaaaacaagttcagggcc	atg	taa
100386	3AORF008	3	2217623630	484	aatgatttagggtaggtgttgacca	atg	tga
100387	3AORF009	1	4072642093	455	gtaaatacttttataagaatggtag	gtg	taa
100388	3AORF010	3	1349114738	415	gaggcggactaacgctacagtaaaa	att	taa
100389	3AORF011	2	20393277	412	attaaagacataatgcgttaaggag	gtg	taa
100390	3AORF012	2	40015209	402	aaaaaagagaaaaaattaaacgcga	atg	taa
100391	3AORF013	1	3037931545	388	attttatgaatgcgagaataaatgc	atg	taa
100392	3AORF014	2	1473815562	274	attatatgggaggtttgactaatta	atg	tag
100393	3AORF015	3	32494034	261	cttgaattaagaaaatctttgaaag	gtg	tag
100394	3AORF016	-2	2558726273	228	aagaagctaagaaaaaaataaaaat	atg	tga
100395	3AORF017	3	67297370	213	ttaatttttaaggaggaaataagca	atg	taa
100396	3AORF018	3	2454025154	204	aataaaataaaaagtaggtgataag	atg	taa
100397	3AORF019	2	3156532128	187	ctataaaaattaaaaaggacggtat	ata	taa
100398	3AORF020	3	3615036713	187	gcagtaggaattatgacgggtcaag	ttg	taa
100399	3AORF021	2	2401124535	174	gtaataaaatttataaagaaaggaa	atg	tga
100400	3AORF022	-2	1242312938	171	taaagtaccagtagacaatgtaggt	att	tga
100401	3AORF023	1	74627917	151	aaaataaatcaaaggagaataattt	atg	taa
100402	3AORF024	1	2673127174	147	actaaataaaataaggaggacact	atg	tga
100403	3AORF025	1	4210642543	145	taagcataagtaatggaggtataag	atg	taa
100404	3AORF026	2	3525535671	138	aagcaactaactttattttaaggag	ata	taa
100405	3AORF027	2	58886298	136	atattggctataatacagtggtttt	atc	taa
100406	3AORF028	-3	2784528255	136	ccttttaagatgtttatgatccttt	ctg	taa
100407	3AORF029	3	3434434748	134	ttaaggttttagatttagaggtgga	atg	taa
100408	3AORF030	2	62996694	131	tataaaaaaggagttggccagataa	atg	tag
100409	3AORF031	1	2083321225	130	ttaacaaaattataggagtgagaaa	ata	taa
100410	3AORF032	-2	3998440361	125	aaatagctgttagagggttacccct	ata	tag
100411	3AORF033	1	79578325	122	gaatatctgcgtcttttttatttga	ata	taa
100412	3AORF034	-2	2850628871	121	gttatcaacctaaggaggtgataac	atg	tag
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100414	3AORF036	2	3002030382	120	accaattttaaggaggagttaatca	atg	tga
100415	3AORF037	2	2181822165	115	aagtgtaagtaatagttaagagtca	gtg	tag
100416	3AORF038	-2	4200342347	114	gtactcactttcaactgcttcaacc	atc	tga
100417	3AORF039	2	2138621727 .	113	tccagaaaatctagagtcataggtt	ata	taa
100418	3AORF040	-3	2965429995	113	ttgattaactcctccttaaaattgg	ttg	taa
100419	3AORF041	-1	43334671	112	tactaaatctacatctgatccatga	att	tga
100420	3AORF042	3	55685900	110	taaaaaagtggtaggtgattttaa	atg	tga
100421	3AORF043	1	2569026019	109	taccaaattaatatagtcttcgcat	ata	tag
100422	3AORF044	3	2967630005	109	gtcttaaataattatataaggagtt	att	taa
100423	3AORF045	3	30353	107	cgctagcaacgcggataaattttc	atg	taa
100424	3AORF046	3	2789428214	106	aagatattgaaaagctaatttcccc	ata	tga
100425	3AORF047	-2	1190712227	106	ttcgccgccaaaatgattagcattt	ctg	tga
100426	3AORF048	-3	4034340663	106	ccataacacatacactgtatgatct	ctg	taa
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WO 00/32825

### PCT/IB99/02040

189

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### Table 9

Bacteriophage 96, complete genome sequence

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193

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tttaaatagc cgttttagca atatgattct aggccataac ggcgacggta tcaatgaagt 32201 aaaagacgcg cgtattgata atacaggtta tggtcataag acattgcaag atcgtttgta tcatgattat 32271 teaacactag atgettteac taaaaaggtt gagaaagctg tagatgaaca etataaagaa tategagega cagaataceg attegaacca aaagagcaag aaceggaatt tateactgat ttategecat atacaaatge 32341 32411 agtaatgcaa tcattttggg tagaccctag aacgaaaatt atttatatga cgcaagctcg tccaggtaat cattacatgt tatctagatt gaagcccaac ggacaattta ttgatagatt gcttgttaaa aacggcggtc 32481 32551 acggtacaca caatgcgtat agatacattg atggagaatt atggatttat tcagctgtat tggacagtaa caaaaacaac aagtttgtac gtttccaata tagaactgga gaaataactt atggtaatga aatgcaagat 32621 32691 gtcatgccga atatattaa cgacagatat acgtcagcga tttataatcc tatagaaaat ttaatgattt 32761 tcagacgtga atataaagct tctgaaagac aagctaagaa ttcattgaat ttcattgaag taagaagtgc 32831 tgacgatatt gataaaggta tagacaaagt attgtatcaa atggatatac ctatggaata cacttcagat acacaaccta tgcaaggtat cacttatgat gcaggtatct tatattggta tacaggtgat tcgaatacag 32901 32971 ccaaccctaa ctacttacaa ggtttcgata taaaaacaaa agaattgtta tttaaacgac gtatcgatat 33041 tggcggtgtg aataataact ttaaaggaga cttccaagaa gctgagggtc tagatatgta ttacgatcta 33111 gaaacaggac gtaaagcact tttaataggg gtaactattg gacctggtaa taacagacat cactcaattt attotatcgg ccaaagaggt gttaaccaat tcttaaaaaa cattgcacct caagtatcga tgactgattc 33181 33251 aggtggacgt gttaaaccgt taccaataca gaacccagca tatctaagtg atattacgga agttggtcat tactatatct atacgcaaga cacacaaaat gcattagatt tcccgttacc gaaagcgttt agagatgcag 33321 33391 ggtggttctt ggatgtactg cctggacact ataatggtgc tctaagacaa gtacttacca gaaacagcac aggtagaaat atgcttaaat tcgaacgtgt cattgacatt ttcaataaga aaaacaacgg agcatggaat 33461 33531 ttottgtccgc aaaacgccgg ttattgggaa catatcccta agagtattac aaaattatca gatttaaaaa tcgttggttt agatttctat atcactactg aagaatcaaa acgatttact gatttccta aagactttaa 33601 33671 aggtattgca ggttggatat tagaagtaaa atcgaataca ccaggtaaca caacacaagt attaagacgt 33741 aataacttcc cgtctgcaca tcaattttta gttagaaact ttggtactgg tggcgttggt aaatggagtt 33811 tattcgaagg 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agegeageta acacatetag gattatteat attactaatg caacagatge gecagaaaag 34721 acggatatag gcacgttaga gaagcctgga Caagatggtg ttgatgacgg ttcttcgttc gatgaatcaa cttatacatc aagcaaatct ggtgtgttag ttgtttatgt tgttgataat aatactgctc gtgcaacatg 34791 34861 gtacccagac gattcaaacg atgagtacac aaaatacaaa atctacggca catggtaccc gttttataaa 34931 aagaatgatg gaaacttaac taagcaattt gttgaagaaa cgtctaaca cgctttaaat caagctaagc agtatgtaga tgataaattc ggaacaacga gctggcaaca acataagatg acagaggcga atggtcaatc 35001 35071 aattcaagtt aacttaaata atgcgcaagg cgatttggga tatttaactg ctggtaatta ctatgcaaca agagtgccgg atttaccagg tagtgttgaa agttatgagg gttatttatc ggtattcgtt aaagacgata 35141 35211 35281 Caaacaagct atttaacttc acgccttata actctaaaaa gatttacaca cgatcaatca caaacggcag acttgagcaa cagtggacag ttcctaatga acataagtca acggtattgt tcgacggtgg agcaaatggt 35351 gtaggtacaa caatcaatct aaccgaacca tacacaaact attctatttt attagtaagt ggaacttatc 35421 caggtggcgt tattgaggga ttcggactaa ccacattacc taatgcaatt caattaagta aagcgaatgt 35491· 35561 agttgactca gacggtaacg gtggcggtat ttatgagtgt ttactatcca aaacaagtag cactacttta agaatcgata acgatgtgta ctttgattta ggtaaaacat caggttctgg agcgaatgcc aacaaagtta 35631 ctataactaa aattatgggg tggaaataat gaaaatcaca gtaaatgata aaaatgaagt tatcggatac 35701 gttaatactg gcggtttacg caatagttta gatgtagacg ataacaatgt gtctatcaaa ttcaaagaag 35841 agttegaace taqaaaqtte qtttteacta acggegaaat taaatacaat agcaattteg aaaaaqaaqa egtacegaat gcatcaaace aacaaagtge gteagattta agtgatgagg aacttegegg aatggttgca 35911 35981 agtatgcaaa tqcaqatqac qcaaqtgaac atqttgacaa tqcaattqac qcaacaaaac qctatqttaa 36051 cacaacagtt gaccgaactg aaaactaaca aaacaaatac tgagggggac gtttaaatga tgaagatgat 36121 ttatccaact tttaaagaca ttaaaacttt ttatgtgtgg ggttgctata aaaatgagca aattaagtgg tacgtagaca tgggtgtaat cgacaaagaa gaatatgcat tgatcactgg tgaaaaatat ccagaggcaa 36191 aagatgaaaa gtcacaggtg taatgcttga ggctttttaa tttaacacaa agtaggtggc gtaatgtttg 36261 gatttaccaa acqqcacqaa catqaatqqc gaattagaag attagaagag aatqataaaa caatqcttaq cacteteaat gagattaaat 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ggtttggtaa atgtaaataa cagttaagag tcagtgcttc ggcactggct ttttattttg attgaaatga 36681 ggtgcataca tgggattacc taacccaaag actagaaagc ctacagctag tgaagtggtg gagtgggcaa 36751 agtoqaatat tqqtaaqaqq attaatatag ataattatog gggcagtcaa tqttgggata cacctaactt 36821 tatttttaaa agatattggg gttttgtaac atggggcaat gctaaggata tggctaatta cagatatcct 36891 aagggtttcc gattctatcg ttattcatct ggatttgtac cggaacctgg agacatcgca gtttggcacc 36961 ctggcaacgg aataggttcg gacggacaca ccgcaatagt agtaggacca tctaataaaa gttatttta 37031 tagcgttgac caaaactggg ttaattctaa tagttggaca ggttctccag gaagattagt aagacacct tatgtaagtg ttacaggct tgttaggcct ccatactcaa aagatactag caaacctagt agtactgata 37101 37171 37241 taaaacagta aaatacactg cttacagcaa tgttttagat aaagaagagc acttcattga tcatatagtt 37311 graatgggtg atgaacgctc agatattcaa ggattatata taaaagaatc aatgcatatg cgttctgtag acgaactgta tacgcaaaga aataagttta taagcgatta tgaaataccg catttatatg tcgatagaga 37381 37451 ggctacatgg cttgctagac caaccaattt tgatgacccg cgtcacccta attggctagt tattgaagta 37521 tgtggtggtc aaacagatag caaacgacaa ttcttattga atcaaataca agcgttaata cgtggtgttt 37591 ggttattgtc agggattgat aaaaacttat ctgaaacgac gttaaaggta gaccctaata tttggcgtag 37661 tatgaaagat ttaattaatt acgacttgat taagcaaggt ataccggata acgcaaagta tgagcaagtt 37731 37801 aaaaagaaaa tgcttgagac atacattaaa cgagatatat tgacacgaga aaatataaaa gaagtaacga 37871 caaaaacaac aataagaatt agtgataaaa catcagttga cagtgcgtcc acacgaggcc ctactccatc agacgaaaaa ccaagcatcg ttactgaaac aagtccattc acattccagc aagcactgga tagacaaatg tctaggggta acccgaaaaa atctcataca tggggctggg ctaatgcaac acgagcacaa acgagctcgg 37941 38011 caatgaatgt taagcgaata tgggaaagta acacgcaatg ctatcaaatg cttaatttag gcaagtatca 38081 aggeatttea gttagtgege ttaacaaaat acttaaagga aaaggaacge tegaeggaca aggeaaagca 38151 ttcgcggaag cttgtaagaa aaacaacatt aacgaaattt atttgatcgc gcacgctttc ttagaaagtg 38221 gatacggaac aagtaacttc gctagtggta gatacggtgc atataattac ttcggtattg gtgcattcga 38291 caacgacct gattatgcaa tgacgtttgc taaaaataaa ggttggacat ctccagcaaa agcaatcatg 38361 38431 ggcggtgcta gcttcgtaag aaaggattac atcaataaag gtcaaaacac attgtaccga attagatgga 38501 atoctaagaa tocagotaco caccaatacg ctactgotat agagtggtgo caacatcaag caagtacaat cgctaagtta tataaacaaa tcggcttaaa aggtatctac ttcacaaggg ataaatataa ataaagaggt 38571 gtgtaaatgt acaaaataaa agatgttgaa acgagaataa aaaatgatgg tgttgactta ggtgacattg 38641 38711 gctgtcgatt ttacactgaa gatgaaaata cagcatctat aagaataggt atcaatgaca aacaaggtcg 38781 tategateta aaageacatg gettaacace tagattacat ttgtttatgg aagatggete tatatteaaa aatgageee ttattatega egatgttgta aaagggttee ttaeetacaa aatacetaaa aaggttatea 38851 38921 aacacgctgg ttatgttcgc tgtaagctgt ttttagagaa agaagaagaa aaaatacatg tcgcaaactt ttettteaat ategitgata giggtatiga ateigetgia geaaaagaaa tegatgitaa atiggiagat 38991 39061 gatgetatta egagaatttt aaaagataac gegacagatt tattgagcaa agaetttaaa gagaaaatag ataaagatgt catttettac ategaaaaga atgaaagtag atttaaaggt gegaaaggtg ataaaggega 39131 accgggacaa cctggtgcga aaggtgatac aggtaaaaaa ggagaacaag gcgcacccgg taaaaacggt 39201 actgtagtat caatcaatcc tgacactaaa atgtggcaaa ttgatggtaa agatacagat atcaaagcag 39271 aacctgagtt attggacaaa atcaatatcg caaatgttga agggttagaa gataaattgc aagaagttaa aaaaatcaaa gatacaactc tcaacgactc taaaacgtat acggattcaa aaattgctga actagttgat 39341 39411 agegegetty aatetatgaa tacattaaga gaattageag aageaataca aaacaactet attteagaaa 39481 gtgtattgca acagattggc tcaaaagtta gtacagaaga ttttgaggaa ttcaaacaaa cactaaacga 39551 tttatatgct ccaaaaaatc ataatcatga tgagcggtat gttttgtcat ctcaagcttt tactaaacaa 39621 39691 caagcggata atttatatca actaaaaagc gcatctcaac cgacggttaa aatttggaca ggaacagaaa atgaatataa ctatatatat caaaaagacc ctaatacact ttacttaatt aaggggtgat ttttatggaa 39761 ggtaatttta aaaatgtaaa gaagtttatt tacgaaggtg aagaatatac aaaagtatat gctggaaata 39831 tccaagtatg gaaaaagcct tcatcttttg taataaaacc cttacctaaa aataaatatc cggatagcat 39901 agaagaatca acagcaaaaat ggacaataaa tggagttgaa cctaataaaa gttatcaggt gacaatagaa 39971 aatgtacgta gcggtataat gagggtttcg caaactaatt taggttcaag tgatttagga atatcaggag 40041 tcaatagegg 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caggtgtatt acctaataac gcaacaatca aatatgacgg cgcatattgc atcaatgggt 41861 41931 atagatggat tacttatatt gctaatagtg gacaacgtcg ctatattgcg acaggagagg tagataaagc

45003							
42001			ttggtaagtt		_	_	
42071	cattaattat	agggaatctt	acagttatta	aataactatt	tggatggatg	ttaatattcc	tatacacttt
42141	ttaacattac	tctcaagatt	taaatgtaga	taacaggcag	gtactacggt	acttgcctat	ttttttgtta
42211	taatgtaatt	acattaccag	taaccaatct	ggcttaaaac	cacatttccg	gtagccaatc	cggctatgca
42281	gaggacttac	ttgcgtaaag	tagtaagaag	ctgactgcat	atttaaacca	cccatactag	ttgctgggtg
42351	gttgttttt	atgttatatt	ataaatgatc	aaaccacacc	acctattaat	ttaggagtgt	ggttatttt
42421	tatgcaaaaa	aaacgaaaaa	aagttcataa	aaagtattgc	atatcacgtt	taaccgtgtt	ataataaggt
42491	ataccagttg	agaggaggat	aaaaagtgtt	agaaaatttt	aaaactatag	cagaaatcgc	cttttataca
42561	atgtcagcaa	ttgccatage	gaaaacattg	aaaaaagacg	ataagtaagt	agacaagccc	gaaagggctg
42631	tctatatata	aattctaaca	ctaaaatact	atgaaaacaa	tttacattat	tttaatcatt	cttatttgga
42701	taaacgtgtt	tttaggcaac	gatataagta	aaagtgttgt	tgcactgctt	actactttac	tgcttatcaa
42771	tttatggaag	agggataaaa	atgacagcaa	taaaagaaat	aattgaatca	atagaaaagt	tattcgaaaa
42841	agaaacggga	tataaaattg	ctaaaaattc	cggattacca	tatcaaactg	tgcaagattt	aagaaatgga
42911	aaaacatctt	tatcagatgc	cagatttaga	acgataataa	agttatacga	gtatcaaaga	tcgcttgaaa
42981	acgaagaaga	taaataaaag	gagccaaaaa	tatgtttgtt	acaaaagaag	aatttaaaac	tttgaatgta
43051	aaagaagtat	ttgaatcagg	taaaaacttt	ataaaaatta	cagatggaag	acatgcaata	tattgggtaa
43121	atgatagata	cgtagtactt	gaccataaaa	aaggcgattt	gtacccgcaa	aaagcatacc	caaaatatat
43191	caaaagaaaa	ttagtaagtt	aaataattag	aaaaccacgt	cttaattgac	gtggttattt	tttaggtttg
43261	cgcgtgtcaa	atacgtgtca	atttagttct	atttctttag	ttttcttct	aaacttaatt	gcttgtaaac
43331	cgcatagtta	taggetttte	agctatatac	caagataaga	tttatcccgc	cgtctccata	aaaatatgct
43401	tggaaacctt	gatttaatgg	ggttttaatc	tagcaagtgt	caaatatgtg	tcaagaaaat	aattttctga
43471	cacgttgacc	ttgctcttt	ttatgttcat	caagtaagtg	agagtaggtg	tctaaagtta	tagatatatt
43541			taatatattc			_	_

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Table 10

# Bacteriophage 96 ORFs list

SID	LAN	FRA	POS	a.a.	RBS sequence	STA	STO
100733	960RF001	1	2599929142	1047	ccttgaatcgaaaggaggttagcct	ttg	taa
100734	960RF002	1	3200833906	632	tttttacgactaaaggaggcaacca	atg	taa
100735	960RF003	1	3010931995	628	ttatattttagataaggagtagcct	atg	taa
100736	960RF004	1	3676038634	624	attttgattgaaatgaggtgcatac	atg	taa
100737	960RF005	3	3390335729	608	gtttattcgaaggaaaggtggttga	ata	taa
100738	960RF006	2	4058942043	484	aatgatttagggtaggtgttgacca	atg	tag
100739	960RF007	1	1865220091	479	tatacacacatactaaacctgaacg	att	tga
100740	960RF008	2	896010201	413	tggcagaatttgggggggataacga	atg	tga
100741	960RF009	2	1744718670	407	gacgcaataacggaagtgatcgtca	atg	tga
100742	960RF010	1	3864739819	390	taaatataaataaagaggtgtgtaa	atg	tga
100743	960RF011	-1	1191195	358	gtagetegeetaceettattatttt	ttg	tga
100744	960RF012	2	2004521013	322	tttaatgacaaattacctgacatag	atg	tga
100745	960RF013	3	2915730098	313	acttattataagggaggtttgttag	ttg	taa
100746	960RF014	1	2192522839	304	agaaaataaagtgaggtaataaaat	atg	tag
100747	960RF015	1	58126591	259	atacacggtaaaggtgggagaatag	atg	taa
100748	960RF016	1	78528607	251	aataaaatgttgaaaggagagaaaa	atg	taa
100749	960RF017	3	34444190	248	aaatttaacattaatatcactttaa	gtg	taa
100750	960RF018	-3	2828129000	239	taagctatgttgaacatcgctagtc	atg	tga
100751	960RF019	3	71887859	223	tttaccgttctaggacgtggtttaa	atg	taa
100752	960RF020	3	2132421908	194	gaagggcaaaaaggagttttgatat	atg	taa
100753	960RF021	3	66127175	187	attaaaaattaattaaaaggacggt	ata	tag
100754	960RF022	2	2453625093	185	aaagaaaaacgaaggagtgtattaa	atg	taa
100755	960RF023	1	52755811	178	catgaaatggtaggaggtatgaaaa	gtg	tag
100756	960RF024	3	1448115014	177	taaaacgataggagataacgaataa	atg	taa
100757	960RF025	2	2515725666	169_	ataaaaaaattgaaaagaggtatat	att	taa
100758	960RF026	-3	1508415590	168	tcattcttaacatagcccttaattc	atg	tga
100759	960RF027	-1	12291732	167	aatagcaaataaaggagtgtaaaac	atg	taa
100760	960RF028	1	1696017454	164	aaggcgtgtgatacagtgaaaacaa	ttg	taa
100761	960RF029	-1	17362227	163	tatgagaaaaggagtcatataaaag	atg	taa
100762	960RF030	1	2553125995	154	ttttcaagagggagagtcgctcgta	ctg	tag
100763	960RF031	2	2363324097	154	tttagtattgaaggtgattctgtag	atc	tag
100764	960RF032	-2	22482706	152	ataagacaccaaaggggtttggcgc	atg	tga
100765	960RF033	-3	3914739605	152	agcatataaatcgtttagtgtttgt	ttg	taa
100766	960RF034	2	1318113615	144	tagaagtcgaaaaagtggaggcaat	ata	taa
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101114	960RF382	-3	3383133935	34	atgttgtttgtaactcgattaagtt	ttg	taa
101115	960RF383	-3	3368733791	34		atc	tga
101116	960RF384	-3	1353013634	34	gttattacgtcttaatacttgtgtt	gtg	tag
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101118	960RF386	1	1225612357	33	tttgattgattgttctagttaagaa	att	taa
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101125	960RF393	3	3928739388	1	aaaaacggtactgtagtatcaatca	atc	tag
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101127	960RF395	-1	1526615367	33	tcgataatttgtatagcttgtttta	atg	tag
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101129	960RF397		1612316224	33	ttatgtgtgcctatcatattaacaa	ttg	tag
101130	960RF398	-2 -2	1364813749	33	tctttaactgaatgttgaatagcat	ttg	tag
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101134	960RF402	-3	3997840079	33	atattcctaaatcacttgaacctaa	att	tga
	960RF403	-3	3860738708	33	atcttcagtgtaaaatcgacagcca	atg	tag
101136	960RF404	-3	2128821389	33	cagacaccgtcttaagtccctttag	ata	taa

WO 00/32825 PCT/IB99/02040

205

#### Table 11

### SEQUENCE INFORMATION FOR PHAGES MATCHING WITH TABLE 1

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Bacteriophage PM2 nuclease cleavage site
  gi|166145|gb|M32695|BM2NCS [166145]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 nucleotide neighbor)
M32693
  Bacteriophage PM2 Hind III fragment 4
  gi|166144|gb|M32693|BM24HIND3 [166144]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 nucleotide neighbor)
M32693
  Bacteriophage PM2 Hind III fragment 4
  gi|166144|gb|M32693|BM24HIND3 [166144]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 nucleotide neighbor)
M32694
  Bacteriophage PM2 Hind III fragment 3
  gi|166143|gb|M32694|BM23HIND3 [166143]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 MEDLINE link)
  Bacteriophage PM2 structural protein gene containing purine/pyrimidine rich
  regions and anti-Z-DNA-IgG binding regions, complete cds
  gij289360|gb|M26134|BM2PROTTV [289360]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 protein link)
J02452
  bacteriophage fi 3'-terminal region ma
  gi|215409|gb|J02452|PFITR3 [215409]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 MEDLINE link)
AF020798
  Bacteriophage Chp1 genome DNA, complete sequence
  gi|217761|dbj|D00624|BCP1 [217761]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 12 protein links, or 1 genome link)
X72793
  Clostridium botulinum C phage BONT/C1, ANTP-139, ANTP-33, ANTP-17, ANTP-70
  genes and ORF-22
  gi|516171|emb|X72793|CBCBONT [516171]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, I MEDLINE link, 6 protein links, or 4 nucleotide neighbors)
X51464
  Clostridium botulinum D Phage C3 gene for excenzyme C3
  gi|14907|emb|X51464|CBDPE3 [14907]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 2 nucleotide neighbors)
  Bacteriophage c-st (from C. botulinum) C1-tox gene for botulinum C1 neurotoxin
  gi|217780|dbj|D90210|CSTC1TOX [217780]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 protein link)
```

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S49407
   type D neurotoxin [bacteriophage d-16 phi, host = C. botulinum, type D, CB16, Genomic, 4087 nt]
   gi|260238|gb|S49407|S49407 [260238]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 1 protein link)
   Bacteriophage phi29 temperature sensitive mutant TS2(98) DNA polymerase gene
   gi|15733|emb|X53370|POTS298 [15733]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 7 nucleotide neighbors)
  Bacteriophage phi29 temperature sensitive mutant TS2(24) DNA polymerase gene
  gi|15731|emb|X53371|POTS224 [15731]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 7 nucleotide neighbors)
X05973
  Bacteriophage phi29 prohead RNA
  gi|15680|emb|X05973|POP29PRO [15680]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE links, or 4 nucleotide neighbors)
V01155
  Left end of bacteriophage phi-29 coding for 15 potential proteins Among
  these are the terminal protein and the proteins encoded by the genes 1, 2 (sus), 3, and (probably) 4
  gi|15659|emb|V01155|POP29B [15659]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 16 protein links, or 16 nucleotide neighbors)
X73097
  Bacteriophage phi-29 left origin of replication
  gi|312194|emb|X73097|BP29ORIL [312194]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 5 nucleotide neighbors)
M14430
  Bacteriophage phi-29 gene-17 gene, complete cds
  gi|215321|gb|M14430|P29G17A [215321]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 6 protein links, or 8 nucleotide neighbors)
M14431
  Bacteriophage phi-29 gene-16 gene, complete cds
  gi|215319|gb|M14431|P29G16A [215319]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 7 nucleotide neighbors)
  Bacteriophage phi-29 DNA, 3' end
  gi|215343|gb|M20693|P29REPINB [215343]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 4 nucleotide neighbors)
M21016
  Bacteriophage phi-29 DNA, 5' end
  gi|215342|gb|M21016|P29REPINA [215342]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 nucleotide neighbor)
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M12456

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Bacteriophage phi-29 genes 9, 10 and 11 encoding p9 tail, incomplete, p10
   connector, complete, and pll lower collar, incomplete, respectively
   gi|215338|gb|M12456|P29P9 [215338]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 3 protein links, or 2 nucleotide neighbors.)
 M14782
   Bacillus phage phi-29 head morphogenesis, major head protein, head fiber
   protein, tail protein, upper collar protein, lower collar protein, pre-neck-
   appendage protein, morphogenesis(13), lysis, morphogenesis(15), encapsidation genes, complete cds
   gi|215323|gb|M14782|P29LATE2 [215323]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 11 protein links, or 11 nucleotide neighbors)
 M26968
   Bacteriophage phi-29 (from Bacillus subtilis) proteins p1 delta-1 genes, complete cds, and the sus1(629) mutation
   gi|341558|gb|M26968|P29P1D1A [341558]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)
 J02448
   Bacteriophage fl, complete genome
   gi|166201|gb|J02448|F1CCG [166201]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 10 protein links, 205 nucleotide neighbors.
  or I genome link)
  Bacteriophage f2 coat protein gene, partial eds
  gi|166228|gb|M24832|F2CRNACA [166228]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 4 nucleotide neighbors)
J02451
  Bacteriophage fd, strain 478, complete genome
  gi|215394|gb|J02451|PFDCG [215394]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 5 MEDLINE links, 10 protein links, 204 nucleotide neighbors,
  or I genome link)
M34834
  Bacteriophage fr replicase gene, 5' end
  gi|166139|gb|M34834|BFRREGRA [166139]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 protein link, or 9 nucleotide neighbors)
M38325
  Bacteriophage fr replicase gene, 5' end
  gi|166137|gb|M38325|BFRREGR [166137]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 protein link, or 9 nucleotide neighbors)
M35063
  Bacteriophage fr coat protein replicase cistron (R region) RNA
  gi|166134|gb|M35063|BFRRCRRA [166134]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 protein link, or 3 nucleotide neighbors)
S66567
  alpha-atrial natriuretic factor/coat protein=fusion polypeptide [human,
  bacteriophage fr, expression vector pFAN15, PlasmidSyntheticRecombinant, 510 nt]
  gi|435742|gb|S66567|S66567 [435742]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 15 nucleotide neighbors)
                                                                                                            ______
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### X15031 Bacteriophage fr RNA genome gi|15071|emb|X15031|LEBFRX [15071] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 4 protein links, 9 nucleotide neighbors, or I genome link) U51233 Mus musculus neutralizing anti-RNA-bacteriophage fr immunoglobulin variable region light chain (IgM) mRNA, partial cds gi|1277150|gb|U51233|MMU51233 [1277150] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 protein link, or 1669 nucleotide neighbors) U51232 Mus musculus neutralizing anti-RNA-bacteriophage fr immunoglobulin variable region heavy chain (lgM) mRNA, partial cds gi|1277148|gb|U51232|MMU51232 [1277148] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 protein link, or 1073 nucleotide neighbors) Bacteriophage If1, complete genome gi|3676280|gb|U02303|B2U02303 [3676280] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 10 protein links, or 1 genome link) V00604 Phage M13 genome gi|14959|emb|V00604|INM13X [14959] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 10 protein links, or 205 nucleotide neighbors) A32252 -Synthetic bacteriophage M13 protein III probe gi|1567340|emb|A32252|A32252 [1567340] (View GenBank report, FASTA report, ASN.1 report, or Graphical view) A32251 Synthetic bacteriophage M13 protein III probe gi|1567339|emb|A32251|A32251 [1567339] (View GenBank report, FASTA report, ASN. 1 report, or Graphical view) M12465 Bacteriophage M13 mp10 mutations in lac operon gi|215210|gb|M12465|M13LACMUT [215210] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 215 nucleotide neighbors) M24177 Synthetic Bacteriophage M13 (clone M13.SV.B12) SV40 early promoter region DNA gi|209416|gb|M24177|SYNSVB12 [209416] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 1 nucleotide neighbor) Synthetic Bacteriophage M13 (clone M13.SV.B11) SV40 early promoter region DNA gi|209415|gb|M24176|SYNSVB11 [209415] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 1 nucleotide neighbor)

-209

## M24175 Synthetic Bacteriophage M13 (clone M13.SV.8) SV40 early promoter region DNA gi|208806|gb|M24175|SYNM13SV8 [208806] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 242 nucleotide neighbors) M19979 Synthetic hybrids; recombinant DNA from bacteriophage M13 and plasmid pHV33 gi|207813|gb|M19979|SYN33M13M (207813) (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 617 nucleotide neighbors) M19565 Synthetic hybrids; recombinant DNA from bacteriophage M13 and plasmid pHV33 gi|207808|gb|M19565|SYN33M13H [207808] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 567 nucleotide neighbors) M19564 Synthetic hybrids; recombinant DNA from bacteriophage M13 and plasmid pHV33 gi|207807|gb|M19564|SYN33M13G [207807] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 12 nucleotide neighbors) M19563 Synthetic hybrids; recombinant DNA from bacteriophage M13 and plasmid pHV33 gi|207806|gb|M19563|SYN33M13F [207806] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 262 nucleotide neighbors) M19561 Synthetic hybrids; recombinant DNA from bacteriophage M13 and plasmid pHV33 gi|207804|gb|M19561|SYN33M13D [207804] (View GenBank report, FASTA report, ASN.1 report, Graphical view, I MEDLINE link, or 27 nucleotide neighbors) M19560 Synthetic hybrids; recombinant DNA from bacteriophage M13 and plasmid pHV33 gi|207803|gb|M19560|SYN33M13C [207803] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, or 1 MEDLINE link) M19559 Synthetic hybrids; recombinant DNA from bacteriophage M13 and plasmid pHV33 gi|207802|gb|M19559|SYN33M13B [207802] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 227 nucleotide neighbors) M10568 Bacteriophage M13 replicative form II, replication origin, specific nick location gi|215220|gb|M10568|M13ORIB [215220] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 650 nucleotide neighbors) M10910 Bacteriophage M13 gene II regulatory region and M13sj1 mutant gi|215209|gb|M10910|M13IIREG [215209] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 72 nucleotide neighbors) Bacteriophage M13 HaeIII restriction fragment DNA gi|215208|gb|M38295|M13HAEIII [215208] (View GenBank report, FASTA report, ASN.1 report, Graphical view, or 67 nucleotide neighbors)

WO 00/32825 PCT/IB99/02040

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-210
 E02067
    DNA encoding a part of Bacteriophage M13 tg 127
    gi|2170311|dbj|E02067|E02067 [2170311]
    (View GenBank report, FASTA report, ASN.1 report, or Graphical view)
 J02467
    Bacteriophage MS2, complete genome
    gi|215232|gb|J02467|MS2CG [215232]
    (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 8 MEDLINE links, 4 protein links, 20 nucleotide neighbors,
   or 1 genome link)
 AJ004950
   Bacteriophage PI ban gene
   gi|3688226|emb|AJ011592|BP1011592 [3688226]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, or 1 protein link)
 U88974
   Bacteriophage P1 structural lytic transgiycosylase (orf47), pep44b (orf44b),
   pep44a (orf44a), and pep43 (orf43) genes, complete cds; and pep42 (orf42) gene, partial cds
   gi|2661099|gb|AF035607|AF035607 [2661099]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 5 protein links, or 1 nucleotide neighbor)
 AJ000741
   Bacteriophage Pl darA operon
   gi|2462938|emb|AJ000741|BPAJ7641 [2462938]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 10 protein links, or 31 nucleotide neighbors
X01828
   Bacteriophage P1 recombinase gene cin
   gi|15133|emb|X01828|MYP1CIN [15133]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 3 nucleonide neighbors)
X98146
   Bacteriophage P1 DNA sequence around the Op88 operator
   gi|1359513|emb|X98146|BP10P880P [1359513]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, or 1 nucleotide neighbor)
S61175
  immI operon: icd=cell division repressor, ant1=antirepressor {promoters
  P51a, P51b) [bacteriophage P1, Genomic, 728 nt]
  gi|385908|gb|S61175|S61175 [385908]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 3 nucleotide neighbors)
X87824
  Bacteriophage P1 gene 26
  gi|861164|emb|X87824|XXBP1G26 [861164]
  (View GenBank report,FASTA report,ASN.1 report,Graphical view, or 1 protein link)
X15638
  Phage P1 DNA for lytic replicon containing promoter P53 and two open reading frames
  gi|15735|emb|X15638|PP1LREP[15735]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 3 protein links, or 24 nucleotide neighbors
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## X17512 Bacteriophage P1 DNA for immunity region immI gi|15479|emb|X17512|P1IMMUNIY [15479] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 2 MEDLINE links, or 4 nucleotide neighbors) X16005 Bacteriophage PI cl gene for Plc1 repressor protein gi|15477|emb|X16005|P1C1 [15477] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 3 nucleotide neighbors) X03453 Bacteriophage P1 cre gene for recombinase protein gi|15135|emb|X03453|MYP1CRE [15135] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 12 nucleotide neighbors.) Bacteriophage P1 c1 gene 5'-region gi|15128|emb|X06561|MYP1C1 [15128] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 4 protein links, or 6 nucleotide neighbors) Bacteriophage P1 genome fragment (IS2 insertion spot). This regions contains four unidentified reading frames and is known as insertion hot spot for IS2 insertion sequences gi|15118|emb|V01534|MYOVP1 [15118] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 4 protein links, or 3 nucleotide neighbors) X56951 Bacteriophage Pl gene10 gi|406728|emb|X56951|BPP1GP10 [406728] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 2 MEDLINE links, 3 protein links, or 1 nucleotide neighbor) K02380 Bacteriophage P1 replication region including repA, parA, and parB genes and incA, incB, and incC incompatibility determinants gi|215652|gb|K02380|PP1REP [215652] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 5 MEDLINE links, 4 protein links, or 8 nucleotide neighbors) X87674 Bacteriophage P1 lydA & lydB genes gi|974763|emb|X87674|BACP1LYD [974763] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 2 nucleotide neighbors) X87673 Bacteriophage P1 gene 17 gi|974761|emb|X87673|BACP117 [974761] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 1 nucleotide neighbor) M16618 Bacteriophage P1 c1 repressor binding sites gi|215600|gb|M16618|PP1C1 [215600] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 3 nucleotide neighbors)

SEG PPICIN

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Bacteriophage P1 cin gene encoding recombinase, cixL recombination site, and 5' end of C invertible element
     gi|215607|gb||SEG_PP1CIN [215607]
    (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 4 nucleotide neighbors)
  K03173
    Bacteriophage P1 C invertible element, right end, and cixR recombination site
    gi|215606|gb|K03173|PP1CIN2 [215606]
    (View GenBank report, FASTA report, ASN.1 report, or Graphical view)
  215605
    Bacteriophage P1 cin gene encoding recombinase, cixL recombination site, and 5' end of C invertible element
    gi|215605|1c1|X01828 [215605]
    (View GenBank report, FASTA report, ASN. 1 report, or Graphical view)
 M25470
    Bacteriophage P1 tail fiber protein gene, complete cds
    gi|341349|gb|M25470|PP1TFPR [341349]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 3 protein links, or 3 nucleotide neighbors)
 M34382
   Bacteriophage P1 sim region proteins, complete cds
   gi|215661|gb|M34382|PP1SIM [215661]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 2 protein links)
 M81956
   Bacteriophage P1 R protein (R) gene, complete cds
   gi|215658|gb|M81956|PP1RP [215658]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 4 nucleotide neighbors)
 M37080
   Bacteriophage P1 mini-P1 plasmid origin of replication
   gi|215657|gb|M37080|PP1REPOR [215657]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 46 nucleotide neighbors)
M27041
  Bacteriophage P1 ref gene, complete cds
  gi|215650|gb|M27041|PP1REF [215650]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 1 nucleotide neighbor)
L01408
  Bacteriophage PI partition protein (parB) gene, 3' end .
  gi|215642|gb|L01408|PP1PARB [215642]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 protein link, or 41 nucleotide neighbors)
SEG PPIPAR
  Bacteriophage miniplasmid P1 parA gene, 5' end
  gi|215639|gb||SEG_PP1PAR [215639]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 48 nucleotide neighbors)
M36425
  Bacteriophage miniplasmid P1 parB gene, 3' end
  gi|215638|gb|M36425|PP1PAR2 [215638]
  (View GenBank report, FASTA report, ASN.1 report, or Graphical view)
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WO 00/32825 PCT/IB99/02040

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213
M36424
   Bacteriophage miniplasmid P1 parA gene, 5' end
  gi|215637|gb|M36424|PP1PAR1 [215637]
  (View GenBank report, FASTA report, ASN.1 report, or Graphical view)
   Bacteriophage P1 miniplasmid origin of replication region
   gi|215632|gb|M11129|PP1ORIM [215632]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 43 nucleotide neighbors)
M25414
   Bacteriophage P1 c1 repressor binding site, operator 88 (Op88)
  gi;215631|gb|M25414|PP1OP88A [215631]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 3 nucleotide neighbors)
M25413
  Bacteriophage P1 c1 repressor binding site, operator 68 (Op68)
  gi|215630|gb|M25413|PP1OP68A [215630]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 MEDLINE link)
  Bacteriophage P1 c1 repressor binding site, operator 21 (Op21)
  gi|215629|gb|M25412|PP1OP21A [215629]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 nucleotide neighbor)
M10510
  Bacteriophage P1 recombination site loxR
  gi|215628|gb|M10510|PP1LOXR [215628]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 nucleotide neighbor)
M10287
  Bacteriophage P1 loxP X loxP recombination site
  gi|215627|gb|M10287|PP1LOXPX [215627]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 13 nucleotide neighbors)
M10494
  Bacteriophage P1 recombination site loxP
  gi|215626|gb|M10494|PP1LOXP [215626]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 134 nucleonde neighbors)
  Bacteriophage P1 recombination site loxL
  gi|215625|gb|M10511|PP1LOXL (215625)
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 nucleotide neighbor)
M10512
  Bacteriophage P1 recombination site loxB
  gi|215624|gb|M10512|PP1LOXB [215624]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 MEDLINE link)
M10145
  Bacteriophage P1 genome fragment with recombination site loxP
  gi|215623|gb|M10145|PP1CREX [215623]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 21 nucleotide neighbors)
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M13327
    Bacteriophage P1 Cin recombinase activated cross over site, junction IV, clone pSHI326
     gi|215622|gb|M13327|PP1CN26IV [215622]
    (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 7 nucleotide neighbors)
  M13325
    Bacteriophage P1 Cin recombinase activated cross over site, junction II, clone pSHI326
    gi|215621|gb|M13325|PP1CN26II [215621]
    (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 1401 nucleotide neighbors)
  M13323
    Bacteriophage P1 Cin recombinase activated cross over site, junction IV, clone pSHI325
    gi|215620|gb|M13323|PP1CN25IV [215620]
    (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 7 nucleotide neighbors)
    Bacteriophage P1 Cin recombinase activated cross over site, junction II, clone pSHI325
    gi|215619|gb|M13321|PP1CN25II [215619]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1058 nucleotide neighbors)
   Bacteriophage P1 Cin recombinase activated cross over site, junction I, clone pSHI326
   gi|215618|gb|M13324|PP1CIR26I [215618]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 7 nucleotide neighbors)
 M13319
   Bacteriophage P1 Cin recombinase activated cross over site, right junction, clone pSHI327
   gi|215617|gb|M13319|PP1CIN27R [215617]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 7 nucleotide neighbors)
 M13320
   Bacteriophage P1 Cin recombinase activated cross over site, junction I, clone pSHI325
   gi|215616|gb|M13320|PP1CIN25I [215616]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 7 nucleotide neighbors)
 M13318
  Bacteriophage P1 Cin recombinase activated cross over site, left junction, clone pSHI324
   gi|215615|gb|M13318|PP1CIN24L [215615]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 1370 nucleotide neighbors)
M13317
  Bacteriophage P1 Cin recombinase activated cross over site, right junction, clone pSHI323
  gi|215614|gb|M13317|PP1CIN23M [215614]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 1055 nucleotide neighbors)
M13316
  Bacteriophage P1 Cin recombinase activated cross over site, left junction, clone pSHI323
  gi|215613|gb|M13316|PP1CIN23L [215613]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 7 nucleotide neighbors)
M13315
  Bacteriophage PI Cin recombinase activated cross over site, right junction, clone pSHI322
  gi|215612|gb|M13315|PP1CIN22R [215612]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 7 nucleotide neighbors)
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215
M13314
  Bacteriophage P1 Cin recombinase activated cross over site, left junction, clone pSHI322
  gi|215611|gb|M13314|PP1CIN22L [215611]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1401 nucleotide neighbors)
M13313
  Bacteriophage P1 Cin recombinase activated cross over site, right junction, clone pSHI321
  gi|215610|gb|M13313|PP1CIN21R [215610]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 7 nucleotide neighbors)
M13312
  Bacteriophage P1 Cin recombinase activated cross over site, left junction, clone pSHI321
  gi|215609|gb|M13312|PP1CIN21L [215609]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1058 nucleotide neighbors)
  Bacteriophage P1 c4 repressor gene, complete cds
  gi|215603|gb|M16568|PP1C4 [215603]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 4 nucleotide neighbors)
M13326
  Bacteriophage P1 Cin recombinase activated cross over site, junction III, clone pSHI326
  gi|215602|gb|M13326|PP1C26III [215602]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1192 nucleotide neighbors)
M13322
  Bacteriophage P1 Cin recombinase activated cross over site, junction III, clone pSHI325
. gi|215601|gb|M13322|PP1C25III [215601]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1231 nucleotide neighbors)
J05651
  Bacteriophage P1 modulator protein (bof) gene, complete cds
  gi|215598|gb|J05651|PP1BOFY1 [215598]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 3 nucleotide neighbors)
M33224
  Bacteriophage P1 regulatory protein (bof) gene, complete cds
  gi|215596|gb|M33224|PP1BOFFO [215596]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 3 nucleotide neighbors)
M10288
  E.coli/bacteriophage P1 loxR recombination site
  gij146647|gb|M10288|ECOLOXR [146647]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 3 nucleotide neighbors)
M10289
  E.coli/bacteriophage P1 loxL recombination site
  gi|146646|gb|M10289|ECOLOXL [146646]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 2 nucleotide neighbors)
M10290
  E.coli loxB site, which can recombine with bacteriophage P1 loxP site
  gi|146645|gb|M10290|ECOLOXB [146645]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 2 nucleotide neighbors)
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M10287

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Bacteriophage P1 loxP X loxP recombination site
      gi|215627|gb|M10287|PP1LOXPX [215627]
      (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 13 nucleotide neighbors)
   M74046
      Bacteriophage P1 pacA and pacB genes, complete cds
      gi|215634|gb|M74046|PP1PACAB [215634]
     (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 2 protein links)
   M95666
      Bacteriophage P1 gene 10, doc and phd genes, complete cds
     gi|463276|gb|M95666|PP1PHDDOC [463276]
     (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 2 MEDLINE links, 4 protein links, or 1 nucleotide neighbor)
     Bacteriophage Q-beta mutated autonomously replicating sequence MDVI RNA fragment
     gi|556359|gb|M25604|PQBARSMUT [556359]
     (View GenBank report, FASTA report, ASN.1 report, Graphical view, I MEDLINE link, or 8 nucleotide neighbors)
   V00643
     first half of the phage Q-beta gene for coat protein
     gi|15088|emb|V00643|LEQBET [15088]
     (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 4 nucleotide neighbors)
  M25167
     Bacteriophage Q-beta RNA fragment recovered from replicase binding complex
     gi|556362|gb|M25167|PQBREPLICB [556362]
    (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 2 nucleotide neighbors)
  M24876
    Bacteriophage Q-beta replicase RNA, 5' end
    gi|556360|gb|M24876|PQBREPLICA [556360]
    (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 4 nucleotide neighbors)
  M25444
    Synthetic bacteriophage Q-beta DNA
    gi|209118|gb|M25444|SYNPQBTERM [209118]
    (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 8 nucleotide neighbors)
M25463
    Bacteriophage Q-beta self-replicating microvariant (+) RNA
    gi|532489|gb|M25463|PQBMVSRRNA [532489]
    (View GenBank report, FASTA report, ASN. 1 report, Graphical view, or 1 MEDLINE link)
 M25014
    Bateriophage Q-beta RNA replicase gene, 5'end, and maturation protein gene, 3' end
    gi|294316|gb|M25014|PQBREPLC [294316]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 2 nucleotide neighbors)
 M25065
    Bacteriophage Q-beta RNA sequence with putative stem loop
   gi|294315|gb|M25065|PQBLOOP [294315]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 3 nucleotide neighbors)
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M10265

217

# Bacteriophage Q-beta RNA molecule with the ability to replicate extracellularly gi|215726|gb|M10265|PQBRNA [215726] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 8 nucleotide neighbors) M24815 Bacteriophage Q-beta specified replicase subunit RNA, gi|215725|gb|M24815|PQBREPL [215725] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 4 nucleotide neighbors) M25461 Bacteriophage Q-beta plus-strand RNA, 5' terminus gi|215724|gb|M25461|PQBPS5E [215724] (View GenBank report, FASTA report, ASN. 1 report, or Graphical view) M25462 Bacteriophage Q-beta plus-strand RNA, 3' terminus gi|215723|gb|M25462|PQBPS3E [215723] (View GenBank report, FASTA report, ASN.1 report, Graphical view, or 8 nucleotide neighbors) M24871 Bacteriophage Q-beta nanovariant WSIII RNA gi|215722|gb|M24871|PQBNVWSIC [215722] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 2 nucleotide neighbors) M24870 Bacteriophage Q-beta nanovariant WSII RNA gi|215721|gb|M24870|PQBNVWSIB [215721] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 2 nucleotide neighbors) Bacteriophage Q-beta nanovariant WSI RNA gi|215720|gb|M24869|PQBNVWSIA [215720] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 2 nucleotide neighbors) M10495 Coliphage Q-beta MDV-1(+) RNA gi|215719|gb|M10495|PQBMDV1A [215719] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 10 nucleotide neighbors) . J02484 bacteriophage qbeta coat protein cistron first half gi|215717|gb|J02484|PQBCP5 [215717] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 4 nucleotide neighbors) M57754 Bacteriophage Q-beta minus strand RNA, 5' terminus gi|215716|gb|M57754|PQBBMS5E [215716] (View GenBank report, FASTA report, ASN.1 report, Graphical view, or 8 nucleotide neighbors) M24297 Bacteriophage Q-beta 5'-terminal region of the minus strand gi|215715|gb|M24297|PQB5END [215715] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 8 nucleotide neighbors)

WO 00/32825 PCT/IB99/02040

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M10695
                                                         218
   Bacteriophage Q-beta, MDV-1 RNA
   gi|215714|gb|M10695|PQB1IR [215714]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE links, or 12 nucleotide neighbors)
 M24827
   Bacteriophage R17 A protein gene, 5' end
   gi|216078|gb|M24827|R17RNACIS [216078]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 5 nucleotide neighbors)
 M24829
   Bacteriophage R17 coat protein gene, 5' end
   gi|216075|gb|M24829|R17CP5 [216075]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 5 nucleotide neighbors)
 J02488
   bacteriophage r17 ma synthetase initiation site
   gi|216080|gb|J02488|R17RNASYN [216080]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 3 MEDLINE links, 2 protein links, or 6 nucleotide neighbors)
 J02487
  bacteriophage r17 coat protein initiation site
  gi|216073|gb|J02487|R17COATP [216073]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, or 1 MEDLINE link)
  bacteriophage r17 a protein initiation site
  gi|216071|gb|J02486|R17APROT [216071]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 MEDLINE link)
M24826
  Bacteriophage R17 coat protein RNA fragment
  gi|216077|gb|M24826|R17CPRAA [216077]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 7 nucleotide neighbors)
M24296
  Bacteriophage R17 3'-terminal fragment A RNA
  gi|216070|gb|M24296|R173TFA [216070]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 9 nucleotide neighbors)
1TFN
  structure refinement for a 24-nucleotide ma hairpin, nmr, minimized average
  structure ribonucleic acid, hairpin, bacteriophage r17 mol_id: 1; molecule: r17c; chain: null; engineered: yes
  gi|1942336|pdb|1TFN| [1942336]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 structure link)
IRPEA
  ma (5'-d(gpgpgpapcpupgpapcpgpapupcpapcpgp срардрирсрирари-3') (24-mer гла
  hairpin coat protein binding site for bacteriophage r17) (nmr, minimized average structure)
  gi|1421020|pdb|1RHT| [1421020]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 structure link)
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M14428
    Bacteriophage S13 circular DNA, complete genome
    gi|216089|gb|M14428|S13CG [216089]
    (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 2 MEDLINE links, 12 protein links, 26 nucleotide neighbors,
    or I genome link)
 J05393
   Bacteriophage T1 DNA N-6-adenine-methyltransferase (M.T1) gene, complete cds
   gi|166163|gb|J05393|BT1NAMTA [166163]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 2 protein links)
 L46845
   Bacteriophage T2 frd3, frd2 genes, complete cds
   gi|951387|gb|L46845|PT2FRD32G [951387]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 protein links, or 17 nucleotide neighbors)
 L43611
   Bacteriophage T2 fibritin (wac) gene, complete cds
   gi|903869|gb|L43611|PT2WAC [903869]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 protein link, or 4 nucleotide neighbors)
 M24812
   Bacteriophage T2 secondary structure RNA sequence
   gi|215796|gb|M24812|PT2RNA [215796]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 4 nucleotide neighbors)
M22342
   Bacteriphage T2 DNA-(adenine-N6)methyltransferase (dam) gene, complete cds
   gi|215792|gb|M22342|PT2DAM [215792]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 2 nucleotide neighbors)
S57515
  orf 61.2 {intergenic region between 41 and 61} [bacteriophage T2, Genomic, 323 nt]
  gi|298524|gb|S57515|S57515 [298524]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 1 protein link)
X05312
  Bacteriophage T2 gene 38 for receptor recognizing protein
  gi|15197|emb|X05312|MYT2G38 [15197]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 protein link)
  Bacteriophage T2 gene 37 for receptor recognizing protein
  gi|15195|emb|X04442|MYT2G37 [15195]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 protein link)
  Bacteriophage T2 gene 32 mRNA for single-stranded DNA binding protein
  gi|15192|emb|X12460|MYT2G32 [15192]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 14 nucleotide neighbors)
X57797
  Bacteriophage T2 gene for gp12
  gi|14875|emb|X56555|BT2GP12 [14875]
 (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 protein link, or 2 nucleotide neighbors)
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X01755
    Bacteriophage T2 tail fiber gene 36
    gi|15189|emb|X01755|MYT2F36 [15189]
    (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)
  M14784
    Bacteriophage T3 strain amNG220B right end, tail fiber protein, lysis protein and DNA packaging proteins, complete cds
    gi|215810|gb|M14784|PT3RE [215810]
    (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 9 protein links, or 10 nucleotide neighbors)
  SEG_PT3RNAPOL
    Bacteriophage T3 RNA polymerase III gene, 5' end
    gi|710559|gb||SEG_PT3RNAPOL [710559]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 2 nucleotide neighbors)
   Bacteriophage T3 RNA polymerase III gene, 3' end
   gi|340722|gb|M22610|PT3RNAPOL2 [340722]
   (View GenBank report, FASTA report, ASN. 1 report, or Graphical view)
 M22609
   Bacteriophage T3 RNA polymerase III gene, 5' end
   gi|340721|gb|M22609|PT3RNAPOL1 [340721]
   (View GenBank report, FASTA report, ASN. 1 report, or Graphical view)
 X05031
   Bacteriophage T3 gene region 1-2.5 with primary origin of replication
   gi|15719|emb|X05031|POT3ORI [15719]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 11 protein links, or 5 nucleotide neighbors)
 X03964
   Bacteriophage T3 early control region pos. 308-810 from genome left end
   gi|15718|emb|X03964|POT3EP [15718]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE links, or 20 nucleotide neighbors)
X17255
   Bacteriophage T3 gene 1 to gene 11
  gi|15682|emb|X17255|POT3111G [15682]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 4 MEDLINE links, 36 protein links, 17 nucleotide neighbors,
   or I genome link)
X15840
  Phage T3 gene 10
  gi|15625|emb|X15840|PODŢ3G10 [15625]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 3 nucleotide neighbors)
  Bacteriophage T3 gene 1 for RNA polymerase
  gi|15561|emb|X02981|PODOT3P [15561]
 (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 3 nucleotide neighbors)
J02503
  bacteriophage 13 5' end, terminally redundant sequence (trs)
  gi|215816|gb|J02503|PT3TRS1 [215816]
  (View GenBank report, FASTA report, ASN.1 report, or Graphical view)
```

## SEG PT3TRS

bacteriophage t3 5' end, terminally redundant sequence (trs)
gi|215818|gb||SEG\_PT3TRS [215818]
(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 1 MEDLINE link)

# J02504

bacteriophage 13 3' end, terminally redundant sequence (115) gi|215817|gb|J02504|PT3TRS2 [215817] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

H YPERLINK http://www.rs.noda.sut.ac.jp/~kunisawa h t t p://www.rs.noda.sut.ac.jp/~kunisawa Bacteriophage T4 genomic database compiled by Arisaka et al.

#### X95646

Bacteriophage T5 DNA for region 60.5%-71% of the T5 genome gi|2791557|emb|AJ001191|BTJ001191 [2791557] (View GenBank report,FASTA report,ASN.1 report,Graphical view,7 MEDLINE links, 12 protein links, or 6 nucleotide neighbors)

#### X56847

Bacteriophage T5 genomic region encoding early genes D10-D15 gij15407|emb|X12930|MYT5D10 [15407] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE/link, 5 protein links, or 4 nucleotide neighbors)

# AF039886

Bacteriophage T5 subclone T5.5.3r5.18r, single pass sequence, genomic survey sequence gi|2811154|gb|AF039886|AF039886 [2811154] (View GenBank report, FASTA report, ASN.1 report, or Graphical view)

#### AF039885

Bacteriophage T5 subclone T5.40f,41f, single pass sequence, genomic survey sequence gi[2811153]gb|AF039885|AF039885 [2811153] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

# AF039884

Bacteriophage T5 subclone T5.26.fr, single pass sequence, genomic survey sequence gi|2811152|gb|AF039884|AF039884 [2811152] (View GenBank report, FASTA report, ASN.1 report, or Graphical view)

# AF039883

Bacteriophage T5 subclone 10-T5.5.7F, single pass sequence, genomic survey sequence gi[2811151]gb]AF039883[AF039883 [2811151] (View GenBank report, FASTA report, ASN. 1 report, or Graphical view)

## AF039882

Bacteriophage T5 subclone 41-T5.5.4BF, single pass sequence, genomic survey sequence gi|2811150|gb|AF039882|AF039882 [2811150] (View GenBank report, FASTA report, ASN.1 report, or Graphical view)

### AF039881

Bacteriophage T5 subclone 39-T5.5.4aF, single pass sequence, genomic survey sequence gi|2811149|gb|AF039881|AF039881 [2811149] (View GenBank report,FASTA report,ASN.1 report,Graphical view, or 1 nucleotide neighbor)

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222

# AF039880

Bacteriophage T5 subclone 19-T5.7.2r, single pass sequence, genomic survey sequence gi|2811148|gb|AF039880|AF039880 [2811148] (View GenBank report, FASTA report, ASN.1 report, or Graphical view)

#### AF039879

Bacteriophage T5 subclone 18-T5.7.2F, single pass sequence, genomic survey sequence gi|2811147|gb|AF039879|AF039879 [2811147] (View GenBank report, FASTA report, ASN.1 report, or Graphical view)

#### AF039878

Bacteriophage T5 subclone 11-T5.5.7R, single pass sequence, genomic survey sequence gi|2811146|gb|AF039878|AF039878 [2811146] (View GenBank report, FASTA report, ASN.1 report, Graphical view, or 2 nucleotide neighbors)

#### AF039877

Bacteriophage T5 subclone T5.4FR, single pass sequence, genomic survey sequence gi|2811145|gb|AF039877|AF039877 [2811145] (View GenBank report, FASTA report, ASN.1 report, or Graphical view)

#### AF039876

Bacteriophage T5 subclone 22-T5.16R, single pass sequence, genomic survey sequence gi|2811144|gb|AF039876|AF039876 [2811144] (View GenBank report, FASTA report, ASN.1 report, or Graphical view)

#### AF039875

Bacteriophage T5 subclone 21-T5.16R, single pass sequence, genomic survey sequence gi|2811143|gb|AF039875|AF039875 [2811143] (View GenBank report, FASTA report, ASN.1 report, or Graphical view)

### AF039874

Bacteriophage T5 subclone 21-T5.16F, single pass sequence, genomic survey sequence gi|2811142|gb|AF039874|AF039874 [2811142] (View GenBank report, FASTA report, ASN.1 report, or Graphical view)

### AF039873

Bacteriophage T5 subclone 09-T5.6F, single pass sequence, genomic survey sequence gi|2811141|gb|AF039873|AF039873 [2811141] (View GenBank report, FASTA report, ASN.1 report, or Graphical view)

### AF039872

Bacteriophage T5 subclone 09-T5.6R, single pass sequence, genomic survey sequence gi|2811140|gb|AF039872|AF039872 [2811140] (View GenBank report, FASTA report, ASN.1 report, Graphical view, or 2 nucleouide neighbors)

## AF039871

Bacteriophage T5 subclone 04-T5.26.R, single pass sequence, genomic survey sequence gi|2811139|gb|AF039871|AF039871 [2811139] (View GenBank report, FASTA report, ASN.1 report, or Graphical view)

# AF039870

Bacteriophage T5 subclone 13-T5.42F, single pass sequence, genomic survey sequence gi|2811138|gb|AF039870|AF039870 [2811138] (View GenBank report, FASTA report, ASN.1 report, or Graphical view)

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X69460
    Bacteriophage T5 ltf gene for L-shaped tail fibers
    gi|15415|emb|X69460|MYT5LTF [15415]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 2 MEDLINE links, 1 protein link, or 4 nucleotide neighbors)
 X03402
   Bacteriophage T5 D15 gene for 5' exonuclease
   gi|15413|emb|X03402|MYT5EXOG [15413]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 2 nucleotide neighbors)
 Z11972
   Bacteriophage T5 tRNA-Tyr, tRNA-Glu, tRNA-Trp, tRNA-Phe, tRNA-Cys and
   tRNA-Asn genes, and ORFs 91aa, 90aa, 42aa and 172aa
   gi|15795|emb|Z11972|T56TRNAG [15795]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 4 protein links, or 3 nucleotide neighbors)
 X03898
   Bacteriophage T5 genes for tRNA-His, -Ser and -Leu
   gi|15786|emb|X03898|STT5RN1 [15786]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, or 2 MEDLINE links)
 X04177
   Bacteriophage T5 gene for transfer RNA-Gln
   gi|15421|emb|X04177|MYT5TRNQ [15421]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 2 nucleotide neighbors)
X03899
   Bacteriophage T5 genes for tRNA-Val, -Lys, -fMet, -Pro and -Ile3
   gi|15787|emb|X03899|STT5RN2 [15787]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, or 1 MEDLINE link)
X03798
  Bacteriophage T5 gene for tRNA-Asp (GUC)
  gil15472|emb|X03798|NCT5TRDG [15472]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 2 nucleotide neighbors)
Y00364
  Bacteriophage T5 tRNA gene cluster (27.8%-22.4%)
  gi|15420|emb|Y00364|MYT5TRN [15420]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 13 nucleotide neighbors)
X03140
  Bacteriophage T5 DNA with rho-dependent transcription terminator (Hind III-P fragment)
  gi|15417|emb|X03140|MYT5RHO [15417]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 2 nucleotide neighbors)
Z35070
  Bacteriophage T6 DNA
  gij535228jembjZ35074jMYEREGBT6 [535228]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 1 protein link)
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AF060870
    Coliphage T6 small subunit distal tail fiber (gene 36) gene, partial cds; and large subunit distal tail fiber (gene 37) and tail fiber
    adhesin (gene 38) genes, complete cds
    gi|3676458|gb|AF052605|AF052605 [3676458]
    (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 3 protein links, or 2 nucleotide neighbors)
 Z35072
    Bacteriophage T6 DNA encoding ORF19.1 gene and g19 gene
    gi|535232|emb|Z35072|MYTAILT6 [535232]
    (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 2 protein links)
 X12488
   Bacteriophage T6 gene 32 mRNA for single-stranded DNA binding protein
   gi|15843|emb|X12488|MYT6G32 [15843]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 14 nucleotide neighbors)
 Z78095
   Bacteriophage T6 DNA (1506 bp)
   gi|1488562|emb|Z78095|BPHZ78095 [1488562]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 protein link, or 4 nucleotide neighbors)
 Z35079
   Bacteriophage T6 DNA for Ip5, Ip6
   gi|535215|emb|Z35079|MY57BT6 [535215]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)
   E.coli bacteriophage T6 gene for beta-glucosyl-HMC-alpha-glucosyl-transferase
  gi|296439|emb|X68725|ECT6 [296439]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 3 protein links, or 1 nucleotide neighbor)
X69894
  Bacteriophage T6 alt gene for ADP-Ribosyltransferase
  gi|15422|emb|X69894|MYT6ADP [15422]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 1 nucleotide neighbor)
L46846
  Bacteriophage T6 frd3, frd2 genes, complete cds
  gi|951390|gb|L46846|PT6FRD32G [951390]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, or 2 protein links)
M27738
  Bacteriophage T6 translational repressor protein (regA), complete cds
  gi|215993|gb|M27738|PT6REGA [215993]
  (View GenBank report, FASTA report, ASN, 1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 5 nucleotide neighbors)
M38465
  Bacteriophage T6 DNA ligase gene, complete cds
  gi|215991|gb|M38465|PT6LIG55 [215991]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 2 nucleotide neighbors)
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V01146
    Genome of bacteriophage T7
    gi|431187|emb|V01146|T7CG [431187]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 13 MEDLINE links, 60 protein links, 105 nucleotide
    neighbors, or 1 genome link)
 X60322
   Bacteriophage alpha3 genes A, B, K, C, D, E, J, F, G, H
    gi|14775|emb|X60322|BACALPHA [14775]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 10 protein links, 22 nucleotide neighbors,
   or I genome link)
 X13332
   Bacteriophage alpha3 DNA for origin of replication
   gi|15093|emb|X13332|MIA3ORPL [15093]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, or 1 MEDLINE link)
 X12611
   Bacteriophage alpha3 gene for protein A part., finger domain
   gi|15092|emb|X12611|MIA3AFIN [15092]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 6 nucleotide neighbors)
X15721
  Bacteriophage alpha3 deletion mutation DNA for the origin region (-ori) of replication
  gi|14774|emb|X15721|BA3DMOR9 [14774]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 11 nucleotide neighbors)
X15720
  Bacteriophage alpha3 deletion mutant DNA for the origin region (-ori) of replication
  gi|14773|emb|X15720|BA3DMOR8 [14773]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 1 nucleotide neighbor)
X15719
  Bacteriophage alpha3 insertion mutant DNA for the origin region (-ori) of replication
  gi|14772|emb|X15719|BA3DMOR7 [14772]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 10 nucleotide neighbors)
  Bacteriophage alpha3 deletion mutation DNA for origin region (-ori) of replication
  gi|14771|emb|X15718|BA3DMOR6 [14771]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 11 nucleonide neighbors)
  Bacteriophage alpha3 deletion mutatnt DNA for origin region (-ori) of replication
  gi|14770|emb|X15717|BA3DMOR5 [14770]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 9 nucleotide neighbors)
X15716
  Bacteriophage alpha3 deletion mutant DNA for origin region (-ori) of replication
 gi|14769|emb|X15716|BA3DMOR4 [14769]
 (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 10 nucleotide neighbors)
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X15715

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Bacteriophage alpha3 deletion mutant DNA for origin region (-ori) of of replication
    gi|14768|emb|X15715|BA3DMOR3 [14768]
    (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 11 nucleonide neighbors)
  X15714
    Bacteriophage alpha3 deletion mutant DNA for origin region (-ori) of replication
    gi|14767|emb|X15714|BA3DMOR2 [14767]
    (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 11 nucleotide neighbors)
 X15713
    Bacteriophage alpha3 deletion mutant DNA for the origin region (-ori) of replication
   gi|14766|emb|X15713|BA3DMOR1 [14766]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 11 nucleotide neighbors)
 X62059
   Bacteriophage alpha3 origin of cDNA synthesis (oriGA)
   gi|14763|emb|X62059|AL3ORIGA [14763]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 13 nucleotide neighbors)
 X62058
   Bacteriophage alpha3 origin of cDNA synthesis (oriAA)
   gi|14762|emb|X62058|AL3ORIAA [14762]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 13 nucleonide neighbors)
 J02444
   Bacteriophage alpha3 origin of DNA replication
   gi|166103|gb|J02444|AL3ORI [166103]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 12 nucleotide neighbors)
   Bacteriophage alpha-3 H protein gene, complete cds
  gi|166101|gb|M25640|AL3HP [166101]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 13 nucleotide neighbors)
  Bacteriophage alpha-3 cleavage site for phage phi-X174 gene A protein
  gi|166099|gb|M10631|AL3CSA [166099]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 3 nucleotide neighbors)
X00774
  Bacteriophage alpha-3 gene J sequence
  gi|15431|emb|X00774|NCBA3J [15431]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 3 protein links, or 2 nucleotide neighbors)
M25640
  Bacteriophage alpha-3 H protein gene, complete cds
  gi|166101|gb|M25640|AL3HP [166101]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 13 nucleotide neighbors)
M10631
  Bacteriophage alpha-3 cleavage site for phage phi-X174 gene A protein
  gi|166099|gb|M10631|AL3CSA [166099]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 3 nucleotide neighbors)
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227 J02459 Bacteriophage lambda, complete genome gi|215104|gb|J02459|LAMCG [215104] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 87 MEDLINE links, 67 protein links, 190 nucleotide neighbors, or 1 genome link) J02482 Bacteriophage phi-X174, complete genome gi|216019|gb|J02482|PX1CG [216019] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 23 MEDLINE links, 11 protein links, 26 nucleotide neighbors, or I genome link) J02454 Bacteriophage G4, complete genome gi|215415|gb|J02454|PG4CG [215415] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 6 MEDLINE links, 11 protein links, 20 nucleotide neighbors. or 1 genome link) X60323 Bacteriophage phiK complete genome gi|1478118|emb|X60323|BPHIKCG [1478118] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 10 protein links, 18 nucleotide neighbors, or 1 genome link) L42820 Bacteriophage BF23 tail protein (hrs) gene, complete cds gi|1048680|gb|L42820|BBFHRS [1048680] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 1 nucleotide neighbor) X54455 Bacteriophage BF23 gene 17 and gene 18 gi|14797|emb|X54455|BF231718G [14797] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 2 protein links, or 2 nucleotide neighbors) M37097 Bacteriophage BF23 DNA, right end of terminal repetition gi|166115|gb|M37097|BBFRIGH [166115] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 2 nucleotide neighbors) M37096 Bacteriophage BF23 DNA, left end of terminal repetition gi|166114|gb|M37096|BBFLEFT [166114] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 nucleotide neighbor) M37095 Bacteriophage BF23 A2-A3 gene, complete cds, and A1 gene, 5' end gi|166110|gb|M37095|BBFA2A3 [166110] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 2 MEDLINE links, 3 protein links, or 1 nucleotide neighbor) AF056281 Bacteriophage BF23 clone bf23.mac5/6.1, genomic survey sequence gi|3090930|gb|AF056281|AF056281 [3090930] (View GenBank report, FASTA report, ASN.1 report, or Graphical view)

#### AF056280

Bacteriophage BF23 clone bf23.mac3, genomic survey sequence gi|3090929|gb|AF056280|AF056280 [3090929] (View GenBank report, FASTA report, ASN. 1 report, or Graphical view)

#### AF056279

Bacteriophage BF23 clone bf23.mac18/21.34, genomic survey sequence gi|3090928|gb|AF056279|AF056279 [3090928] (View GenBank report, FASTA report, ASN.1 report, or Graphical view)

#### AF056278

Bacteriophage BF23 clone bf23.mac16/19.33, genomic survey sequence gi|3090927|gb|AF056278|AF056278 [3090927] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

#### A F056277

Bacteriophage BF23 clone bf23.mac16/19-33, genomic survey sequence gij3090926|gb|AF056277|AF056277 [3090926] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

#### AF056276

Bacteriophage BF23 clone bf23.mac12/9-9, genomic survey sequence gi|3090925|gb|AF056276|AF056276 [3090925] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

# AF056275

Bacteriophage BF23 clone bf23.mac11/14-24, genomic survey sequence gi|3090924|gb|AF056275|AF056275 [3090924] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

### AF056274

Bacteriophage BF23 clone bf23.57r64r, genomic survey sequence gi|3090923|gb|AF056274|AF056274 [3090923] (View GenBank report,FASTA report,ASN.1 report,Graphical view, or 3 nucleotide neighbors )

### AF056273

Bacteriophage BF23 clone bf23.54fr, genomic survey sequence gi|3090922|gb|AF056273|AF056273 [3090922] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

# AF056272

Bacteriophage BF23 clone bf23.47fr.mac10/7, genomic survey sequence gi|3090921|gb|AF056272|AF056272 [3090921] (View GenBank report FASTA report ASN.1 report, or Graphical view)

# AF056271

Bacteriophage BF23 clone bf23.23.66r, genomic survey sequence gi|3090920|gb|AF056271|AF056271 [3090920] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

# AF056270

Bacteriophage BF23 clone bf23.23.64f, genomic survey sequence gi|3090919|gb|AF056270|AF056270 [3090919] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

# AF056269

Bacteriophage BF23 clone bf23.23.60r, genomic survey sequence gi|3090918|gb|AF056269|AF056269 [3090918] (View GenBank report, FASTA report, ASN.1 report, or Graphical view)

#### AF056268

Bacteriophage BF23 clone bf23.23.60f, genomic survey sequence gi|3090917|gb|AF056268|AF056268 [3090917] (View GenBank report,FASTA report,ASN.1 report,Graphical view, or 1 nucleotide neighbor)

## AF056267

Bacteriophage BF23 clone bf23.23.59r, genomic survey sequence gi|3090916|gb|AF056267|AF056267 [3090916] (View GenBank report, FASTA report, ASN.1 report, or Graphical view)

#### AF056266

Bacteriophage BF23 clone bf23.23.59f, genomic survey sequence gi|3090915|gb|AF056266|AF056266 [3090915] (View GenBank report, FASTA report, ASN.1 report, or Graphical view)

#### AF056265

Bacteriophage BF23 clone bf23.23.56r, genomic survey sequence gi|3090914|gb|AF056265|AF056265 [3090914] (View GenBank report, FASTA report, ASN.1 report, or Graphical view)

#### AF056264

Bacteriophage BF23 clone bf23.23.56f, genomic survey sequence gi|3090913|gb|AF056264|AF056264 [3090913] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

#### AF056263

Bacteriophage BF23 clone bf23.23.68f55r, genomic survey sequence gi|3090912|gb|AF056263|AF056263 [3090912] (View GenBank report, FASTA report, ASN.1 report, or Graphical view)

### A F056262

Bacteriophage BF23 clone bf23.23.43fr.66f, genomic survey sequence gi|3090911|gb|AF056262|AF056262 [3090911] (View GenBank report, FASTA report, ASN.1 report, or Graphical view)

### AF056261

Bacteriophage BF23 clone bf23.23.2fr, genomic survey sequence gi|3090910|gb|AF056261|AF056261 [3090910] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

# AF056260

Bacteriophage BF23 clone bf23.23.55.f, genomic survey sequence gi|3090909|gb|AF056260|AF056260 [3090909] (View GenBank report, FASTA report, ASN.1 report, or Graphical view)

# AF056259

Bacteriophage BF23 clone bf23.23.53.r, genomic survey sequence gi|3090908|gb|AF056259|AF056259 [3090908] (View GenBank report, FASTA report, ASN.1 report, or Graphical view)

# AF056258

Bacteriophage BF23 clone bf23.23.53.f, genomic survey sequence gi|3090907|gb|AF056258|AF056258 [3090907] (View GenBank report, FASTA report, ASN.1 report, or Graphical view)

#### AF056257

Bacteriophage BF23 clone bf23.23.52.r, genomic survey sequence gi|3090906|gb|AF056257|AF056257 [3090906] (View GenBank report, FASTA report, ASN.1 report, or Graphical view)

#### AF056256

Bacteriophage BF23 clone bf23.23.52.f, genomic survey sequence gi|3090905|gb|AF056256|AF056256 [3090905] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

#### AF056255

Bacteriophage BF23 clone bf23.23.49.r, genomic survey sequence gi|3090904|gb|AF056255|AF056255 [3090904] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

#### AF056254

Bacteriophage BF23 clone bf23.23.49.f, genomic survey sequence gi|3090903|gb|AF056254|AF056254 [3090903] (View GenBank report, FASTA report, ASN.1 report, or Graphical view)

#### AF056253

Bacteriophage BF23 clone bf23.23.48.r, genomic survey sequence gi|3090902|gb|AF056253|AF056253 [3090902] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

# AF056252

Bacteriophage BF23 clone bf23.23.48.f, genomic survey sequence gij3090901|gb|AF056252|AF056252 [3090901] (View GenBank report, FASTA report, ASN.1 report, or Graphical view)

## AF056251

Bacteriophage BF23 clone bf23.23.44.r, genomic survey sequence gi|3090900|gb|AF056251|AF056251 [3090900] (View GenBank report, FASTA report, ASN.1 report, or Graphical view)

# AF056250

Bacteriophage BF23 clone bf23.23.41.f, genomic survey sequence gi|3090899|gb|AF056250|AF056250 [3090899] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

# AF056249

Bacteriophage BF23 clone bf23.23.22.a.r, genomic survey sequence gi|3090898|gb|AF056249|AF056249 [3090898] (View GenBank report, FASTA report, ASN.1 report, or Graphical view)

# AF056248

Bacteriophage BF23 clone bf23.23.22.a.f, genomic survey sequence gi|3090897|gb|AF056248|AF056248 [3090897] (View GenBank report, FASTA report, ASN.1 report, or Graphical view)

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AF056247
     Bacteriophage BF23 clone bf23.23.68.r, genomic survey sequence
     gi|3090896|gb|AF056247|AF056247 [3090896]
    (View GenBank report, FASTA report, ASN. 1 report, or Graphical view)
  Z50114
    Bacteriophage BF23 DNA for putative tail protein gene
     gi|2464952|emb|Z50114|BF23LATE [2464952]
    (View GenBank report, FASTA report, ASN. 1 report, Graphical view, or 1 protein link)
  D12824
    Bacteriophage BF23 genes for minor tail protein gp24 and major tail protein gp25, complete cds
    gi|520578|dbj|D12824|BBF2TAIL [520578]
    (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 3 nucleotide neighbors)
  Z34953
    Bacteriophage K3 ip9, ip7 and ip8 genes
    gi|535261|emb|Z34953|MYK3IP978 [535261]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 3 protein links, or 1 nucleotide neighbor)
    Bacteriophage K3 DNA for Ip3 and Ip4
   gi|535229|emb|Z35075|MYEORF64K [535229]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 2 protein links)
 X05560
   Bacteriophage K3 gene 38 for receptor recognizing protein
   gi|15112|emb|X05560|MYK3G38 [15112]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 1 protein link)
 X04747
   Bacteriophage K3 gene 37 for receptor recognizing protein
   gi|15110|emb|X04747|MYK3G37 [15110]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 2 nucleotide neighbors)
   Bacteriophage K3 tail fiber gene 36
   gi|15108|emb|X01754|MYK3F36 [15108]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 2 protein links)
  Bacteriophage K3 't' lysis gene, complete cds
  gi|215503|gb|M16812|PK3LYST [215503]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 4 nucleotide neighbors)
L46833
  Bacteriophage K3 frd3, frd2 genes, complete cds
  gi|951377|gb|L46833|PK3FRD32G [951377]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 protein links, or 2 nucleotide neighbors)
L43613
  Bacteriophage K3 fibritin (wac) gene, complete cds
  gi|903861|gb|L43613|PK3WAC [903861]
 (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 protein link, or 4 nucleotide neighbors)
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X01753
      Bacteriophage Ox2 tail fiber gene 36
      gi|15122|emb|X01753|MYOX2F36 [15122]
      (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)
    L43612
      Bacteriophage Ox2 fibritin (wac) gene, complete cds
      gi|903848|gb|L43612|OX2WAC [903848]
      (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 protein link, or 4 nucleotide neighbors)
   Z46880
      Bacteriophage OX2 stp gene
      gi|599663|emb|Z46880|BPOX2STP [599663]
     (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 4 nucleotide neighbors)
   X05675
     Bacteriophage Ox2 gene 38 for receptor-recognizing protein and flanking regions
     gi|15124|emb|X05675|MYOX2G38 [15124]
    (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 3 protein links, or 1 nucleotide neighbor)
  M33533
    Bacteriophage RB18 translational repressor protein (regA) and Orf43.1, complete cds
    gi|216083|gb|M33533|RB18REGA [216083]
    (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 2 nucleotide neighbors)
  AF033329
    Bacteriophage RB18 single-stranded binding protein (gene 32) gene, partial cds, and 5' region
    gi|2645788|gb|AF033329|AF033329 [2645788]
    (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 protein link, or 11 nucleotide neighbors)
  M86231
    Bacteriophage RB69 gene 62, 3'end; RegA (regA) gene, complete cds
    gi|215354|gb|M86231|P6962REGA [215354]
    (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)
  AF033332
   Bacteriophage RB69 single-stranded binding protein (gene 32) gene, partial eds, and 5' region
   gi|2645794|gb|AF033332|AF033332 [2645794]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 protein link, or 12 nucleotide neighbors)
U34036
   Bacteriophage RB69 DNA polymerase (43) gene, complete eds
   gi|1237125|gb|U34036|BRU34036 [1237125]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 1 protein link)
 V01145
   Bacteriophage H1 genome fragment Each Thymine given in this sequence represents a HMU-residue
   (HMU = 5-hydroxymethyluracil)
   gi|15557|emb|V01145|PODOH1 [15557]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 MEDLINE link)
 X05676
   Bacteriophage M1 gene 38 for receptor recognizing protein and flanking regions
   gi|15114|emb|X05676|MYM1G38 [15114]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 3 protein links, or 1 nucleotide neighbor)
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AF034575
     Bacteriophage M1 putative integrase (int) gene, complete cds, and attP region, complete sequence
     gi|2662472|gb|AF034575|AF034575 [2662472]
     (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 protein link)
   AF033321
     Bacteriophage M1 single-stranded binding protein (gene 32) gene, partial cds, and 5' region
     gi|2645772|gb|AF033321|AF033321 [2645772]
    (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 protein link, or 17 nucleotide neighbors)
  X55190
    Bacteriophage Tula 37 and 38 genes for receptor-recognizing proteins 37 and 38 (respectively), partial cds
    gi|14860|emb|X55190|BPTUIA [14860]
    (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 2 nucleotide neighbors)
  AF033334
    Bacteriophage TuIb single-stranded binding protein (gene 32) gene, partial cds, and 5' region
    gi|2645798|gb|AF033334|AF033334 [2645798]
    (View GenBank report, FASTA report, ASN. 1 report, Graphical view, or 5 nucleotide neighbors)
 X55191
    Bacteriophage Tulb 37 gene for receptor-recognizing protein 37 (partial cds), 38 gene for receptor-recognizing protein 38,
    and t gene (partial cds)
    gi|14863|emb|X55191|BPTUIB [14863]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 3 protein links, or 3 nucleotide neighbors)
 X13065
   Bacteriophage phi80 early region
   gil14800|emb|X13065|BP80ER [14800]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 8 protein links, or 6 nucleotide neighbors)
 D00360
   Bacteriophage phi80 cor gene
   gi|217782|dbj|D00360|P8080COR [217782]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 protein link)
X01639
   Bacteriophage phi 80 DNA-fragment with replication origin
   gi|15828|emb|X01639|XXPHI80 [15828]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 25 nucleotide neighbors)
X04051
  Lambdoid bacteriophage phi 80 int-xis region (integrase-excisionase region)
  gi|15770|emb|X04051|STPHI80X [15770]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)
X06751
  Phage Phi80 DNA for major coat protein
  gi|15768|emb|X06751|STPHI80C [15768]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 11 nucleotide neighbors)
X75949
  Bacteriophage phi80 DNA for ORF x171.8 and ORF x171.28'
  gi|458811|emb|X75949|ECORF171B [458811]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 28 nucleotide neighbors)
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L40418
   Bacteriophage phi-80 gene, complete cds
   gi|1019107|gb|L40418|P80A [1019107]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 1 protein link)
 M24831
   Bacteriophage phi-80 Tyr-tRNA gene, 3' end
   gi|215363|gb|M24831|P80TGY [215363]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 43 nucleotide neighbors)
 M10670
   Bacteriophage phi-80 replication origin
   gi|215361|gb|M10670|P80ORI [215361]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 1 nucleotide neighbor)
 M24825
   Bacteriophage phi-80 RNA fragment
   gi|215360|gb|M24825|P80M3A [215360]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 nucleotide neighbor)
 M11919
   Bacteriophage phi-80 cI immunity region encoding the N gene
   gi|215358|gb|M11919|P80CI (215358)
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 2 nucleotide neighbors)
M10891
  Bacteriophage phi-80 attP site DNA
  gi|215357|gb|M10891|P80ATTP [215357]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 nucleotide neighbor)
  Bacteriophage 933J (from E.coli) proviral Shiga-like toxin type 1 subunits A and B genes, complete cds
  gi|215072|gb|M19473|J93SLTI [215072]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE links, 2 protein links, or 20 nucleotide neighbors)
Y10775
  Bacteriophage 933W ileX, stx2A and stx2B genes
  gi|1938206|emb|Y10775|BP933ILEX [1938206]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 protein links, or 36 nucleotide neighbors)
X83722
  Bacteriophage 933W slt-IIB gene
  gi|1490229|emb|X83722|B933WSLT [1490229]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 protein links, or 20 nucleotide neighbors)
X07865
  Bacteriophage 933W slt-II gene for Shiga-like toxin typeII subunit A and B
  gi|14892|emb|X07865|BWSLTII [14892]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 protein links, or 29 nucleotide neighbors)
M16625
  Bacteriophage H19B (from E.coli) sltIA and sltIB genes encoding Shiga-like toxin I subunits A and B, complete cds
  gi|215043|gb|M16625|H19BSLT [215043]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 24 nucleotide neighbors)
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735 M17358 Bacteriophage H19B shiga-like toxin-1 (SLT-1) A and B subunit DNA, complete cds gi|215046|gb|M17358|H19BSLTA [215046] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 20 nucleotide neighbors) 1129728 Bacteriophage N4 single-stranded DNA-binding protein (N4SSB) gene, complete cds gi|939708|gb|U29728|BNU29728 [939708] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 2 MEDLINE links, or 1 protein link) 102580 Bacteriophage PA-2 (E.coli porcine strain isolate) Rz gene, 5'end; ORF2, outer membrane porin protein (lc) and ORF1 genes. complete cds gi|215366|gb|J02580|PA2LC [215366] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 4 protein links, or 4 nucleotide neighbors) U32222 Bacteriophage 186, complete sequence gi|3337249|gb|U32222|B1U32222 [3337249] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 6 MEDLINE links, 46 protein links, or 5 nucleotide neighbors) X51522 Bacteriophage P4 complete DNA genome gi|450916|emb|X51522|MYP4CG [450916] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 3 MEDLINE links, 13 protein links, 6 nucleotide neighbors. or I genome link) Bacteriophage 82 orf33, orf151, orf56, orf96, rus, orf45, and Q genes gi|1051111|emb|X92588|BAC82HOLL [1051111] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 7 protein links, or 1 nucleotide neighbor) J02803 Bacteriophage 82 antitermination protein (Q) gene, complete cds gi|215364|gb|J02803|P82Q [215364] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINElink, or 1 protein link) U02466 Bacteriophage HK022 (cro), (cII) and (O) genes, complete cds, (P) gene, partial cds gi|407285|gb|U02466|BHU02466 [407285] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 5 protein links, or 1 nucleotide neighbor) M26291 Bacteriophage D108 regulatory DNA-binding protein (ner) gene, complete cds gi|166194|gb|M26291|D18NER [166194] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 1 nucleotide neighbor) M11272 Bacteriophage D108 left-end DNA gi|166193|gb|M11272|D18LEDNA [166193] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 2 nucleotide neighbors) M18902 Bacteriophage D108 kil gene encoding a replication protein, 3' end; and containing three ORFs, complete eds gi|166191|gb|M18902|D18KIL [166191] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 3 nucleotide neighbors)

M10191

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Bacteriophage D108, left end with Mu A protein binding sites L1 and L2
    gi|166190|gb|M10191|D18BSL [166190]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 5 nucleotide neighbors)
  J02447
   bacteriophage d108 gene a 5' end
   gi|166189|gb|J02447|D18AAA [166189]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 MEDLINE link)
 V00865
   Bacteriophage D108 fragment from genes A and ner (C-terminus of ner and N-terminus of A)
   gi|15437|emb|V00865|NCD108 [15437]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 2 protein links)
 X01914
   Bacteriophage IKe gene for DNA binding protein
   gi|14957|emb|X01914|INIKEDBP [14957]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 2 nucleotide neighbors)
 AF064539
   Bacteriophage N15, complete genome
   gi|3192683|gb|AF064539|AF064539 [3192683]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE links, 60 protein links, 26 nucleotide neighbors.
   or I genome link )
 U02303
  Bacteriophage If1, complete genome
  gi|3676280|gb|U02303|B2U02303 [3676280]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 10 protein links, or 1 genome link)
AF007792
  Bacteriophage Mu late morphogenetic region
  gij3551775|gb|AF007792|AF007792 [3551775]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 nucleotide neighbor)
U24159
  Bacteriophage HP1 strain HP1c1, complete genome
  gi|1046235|gb|U24159|BHU24159 [1046235]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 6 MEDLINE links, 41 protein links, 8 nucleotide neighbors.
  or I genome link)
Z71579
  Bacteriophage S2 type A 5.6 kb DNA fragment
  gi|1679806|emb|Z71579|BPHS1ADNA [1679806]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 3 MEDLINE links, 9 protein links, or 9 nucleotide neighbors)
X53238
  Klebsiella sp. bacteriophage K11 gene 1 for RNA polymerase
  gi|14984|emb|X53238|KSK11RPO [14984]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 1 nucleotide neighbor)
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X85010

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Bacteriophage A511 ply511 gene
    gi|853748|emb|X85010|BPA511PLY [853748]
    (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 3 protein links, or 1 nucleotide neighbor)
  U29728
    Bacteriophage N4 single-stranded DNA-binding protein (N4SSB) gene, complete cds
    gi|939708|gb|U29728|BNU29728 [939708]
    (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 2 MEDLINE links, or 1 protein link)
    bacteriophage bol 3'-terminal region ma
    gi|166152|gb|J02445|BO1TR3 [166152]
    (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 5 nucleotide neighbors)
  L06183
    Bacteriophage L5 (from Leuconostoc oenos) genome
    gi|289353|gb|L06183|BL5GENM [289353]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, or 1 genome link)
 AF074945
   Mycoplasma arthritidis bacteriophage MAV1, complete genome
   gi|3511243|gb|AF074945|AF074945 [3511243]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 15 protein links, 3 nucleotide neighbors, or 1 genome link)
 L13696
   Bacteriophage L2 (from Mycoplasma), complete genome
   gi|289338|gb|L13696|BL2CG [289338]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 3 MEDLINE links, 14 protein links, or 1 genome link)
 X80191
   Bacteriophage PP7 mRNA for maturation, coat, lysis and replicase proteins
   gi|517237|emb|X80191|BPP7PR [517237]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 4 protein links, or 1 genome link)
 M19377
   Bacteriophage Pf3 from Pseudomonas aeruginosa (New York strain), complete genome
   gi|215380|gb|M19377|PF3COMNY [215380]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 9 protein links, or 5 nucleotide neighbors)
M11912
  Bacteriophage Pf3 from Pseudomonas aeruginosa (Nijmegen strain), complete genome
  gi|215371|gb|M11912|PF3COMN [215371]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 9 protein links, 5 nucleotide neighbors, or 1
  genome link)
V00605
  Bacteriophage Pf1 gene encoding DNA binding protein
  gi|14970|emb|V00605|INOPF1 [14970]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 proteine link, or 1 nucleotide neighbor)
L05626
  Bacteriophage PR4 capsid protein (P6) gene, complete cds
  gi|215735|gb|L05626|PR4P6MAJA [215735]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 1 nucleotide neighbor)
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## D13409

Bacteriophage phiCTX (isolated from Pseudomonas aeruginosa) cosR, attP, int genes gi[217776]dbj[D13409]BPHCOSR [217776]

(View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 3 protein links, or 3 nucleotide neighbors)

# D13408

Bacteriophage phiCTX (isolated from Pseudomonas aeruginosa) cosL, ctx genes gi/217775|dbj/D13408|BPHCOSLCTX [217775]

(View GenBank report, FASTA report, ASN. 1 report, Graphical view, 2 MEDLINE links, or 3 nucleotide neighbors)

#### M24832

Bacteriophage f2 coat protein gene, partial cds gi|166228|gb|M24832|F2CRNACA [166228]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 4 nucleotide neighbors)

#### S72011

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds gi|2618967|gb|AF017629|AF017629 [2618967]

(View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINElink, 2 protein links, or 44 nucleotide neighbors)

#### AF017628

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds gi|2618964|gb|AF017628|AF017628 [2618964]

(View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINElink, 2 protein links, or 44 nucleotide neighbors)

#### AF017627

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds gi|2618961|gb|AF017627|AF017627 [2618961]

(View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINElink, 2 protein links, or 44 nucleotide neighbors)

### AF017626

Bacteriophage 21 isocitrate dehydrogenase (icd) gene, partial cds; and integrase (int) gene, partial cds gil2618958|gb|AF017626|AF017626 [2618958]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 49 nucleotide neighbors)

### AF017625

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds gi|2618955|gb|AF017625|AF017625 [2618955]

(View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINElink, 2 protein links, or 44 nucleotide neighbors)

# AF017624

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int)genes, partial cds gi|2618952|gb|AF017624|AF017624 [2618952]

(View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINElink, 2 protein links, or 44 nucleotide neighbors)

### AF017623

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds gi|2618949|gb|AF017623|AF017623 [2618949]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors)

# AF017622

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds gi[2618946]gb[AF017622|AF017622 [2618946]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors)

239 AF017621 Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds gi|2618943|gb|AF017621|AF017621 [2618943] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors) D26449 Bacteriophage PS17 FI gene for tail sheath protein (gpFI) and FII gene for tail tube protein (gpFII), complete cds gi|452162|dbj|D26449|BPSFIFII [452162] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, or 2 protein links) X87627 Bacteriophage D3112 A and B genes gi|974768|emb|X87627|BPD3112AB [974768] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINElink, 2 protein links, or 1 nucleotide neighbor) U32623 Bacteriophage D3 transcriptional activator CII (cII) gene, complete cds gi|984852|gb|U32623|BDU32623 [984852] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 protein link, or 1 nucleotide neighbor) Bacteriophage phi 11 holin homologue (ORF3) gene, complete cds and peptidoglycan hydrolase (lytA) gene, partial cds gi|511838|gb|L34781|BPHHOLIN [511838] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 4 protein links, or 2 nucleotide neighbors) Bacteriophage P22 (gp10) gene, complete cds, and (gp26) gene, complete cds gi|294053|gb|L14810|P22GP1026X [294053] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 2 nucleotide neighbors) X87420 Bacteriophage ES18 genes 24, c2, cro, c1, 18, and oL and oR operators gi|1143407|emb|X87420|BPES18GEN [1143407] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 5 protein links, or 9 nucleotide neighbors) L42820 Bacteriophage BF23 tail protein (hrs) gene, complete cds gi|1048680|gb|L42820|BBFHRS [1048680] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINElink, 1 protein link, or 1 nucleotide neighbor) X14980 Bacteriophage PRD1 XV gene for protein P15 (lytic enzyme) gi|15802|emb|X14980|TEPRD1XV [15802] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINElink, 1 protein link, or 4 nucleotide neighbors) X06321 Bacteriophage PRD1 gene 8 for DNA terminal protein gi|15800|emb|X06321|TEPRD18 [15800] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 10 nucleotide neighbors) X14336 Filamentous Bacteriophage I2-2 genome gi|14920|emb|X14336|INBI22 [14920] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 9 protein links, 1 nucleotide neighbor, or 1 genome link )

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240
  L05001
    Bacteriophage X glucosyl transferase gene, complete cds
    gi|216044|gb|L05001|PXFCLUSYLT [216044]
    (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 1 protein link)
  M29479
    Bacteriophage p4 sid and psu genes partial cds, and delta gene, complete cds gi|215701|
    gb|M29479|PP4SDP [215701]
    (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 3 protein links, or 4 nucleotide neighbors)
 SEG PP4PSUSID
    Bacteriophage P4 capsid size determination protein (sid) gene, 5' end
    gi|215698|gb||SEG_PP4PSUSID [215698]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)
 M29650
    Bacteriophage P4 polarity suppression protein (psu) gene, complete cds
    gi|215697|gb|M29650|PP4PSUSID2 [215697]
   (View GenBank report, FASTA report, ASN.1 report, or Graphical view)
 M29651
   Bacteriophage P4 capsid size determination protein (sid) gene, 5' end
   gi|215696|gb|M29651|PP4PSUSID1 [215696]
   (View GenBank report, FASTA report, ASN. 1 report, or Graphical view)
 M27748
   Bacteriophage P4 gop, beta, and cII genes, complete cds and int gene, 3' end
   gi|215691|gb|M27748|PP4GOPBC [215691]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 4 protein links, or 1 nucleotide neighbor)
K02750
   Bacteriophage IKe, complete genome
   gij215061|gb|K02750|IKECG [215061]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINElink, 10 protein links, 4 nucleotide neighbors, or 1
   genome link)
L40418
  Bacteriophage phi-80 gene, complete cds
  gi|1019107|gb|L40418|P80A [1019107]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 protein link)
AF032122
  Bacteriophage SfII integrase (int) gene, partial cds; and bactoprenol glucosyl transferase (bgt), and glucosyl transferase II (gttl)
  genes, complete cds
  gi|2465412|gb|AF021347|AF021347 [2465412]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINElink, 4 protein links, or 2 nucleotide neighbors)
M35825
  Bacteriophage SF6 fragment D lysozyme gene, complete cds
  gij216105|gb|M35825|SF6LYZ [216105]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, or 1 protein link)
235479
  Bacteriophage C16 ip1 gene
  gi|534936|emb|Z35479|BC16IP1 [534936]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 2 nucleotide neighbors)
                                                                                                              المحدود
معامل المحادث
المحادث
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X12638

# Bacteriophage 21 DNA for gene 2 gi|296141|emb|X12638|B21GENE2 [296141] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 1 nucleotide neighbor) X02501 Bacteriophage 21 DNA for left end sequence with genes 1 and 2 gi;15825|emb|X02501|XXPHA21 [15825] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 3 nucleotide neighbors) M65239 Bacteriophage 21 lysis genes S, R, and Rz, complete cds gi|215466|gb|M65239|PH2LYSGEN [215466] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 3 protein links, or 1 nucleotide neighbor) M58702 Bacteriophage 21 late gene regulatory region gi|215465|gb|M58702|PH2LATEGE [215465] (View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 MEDLINE link) M81255 Bacteriophage 21 head gene operon gi|215454|gb|M81255|PH2HEADTL [215454] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE links, 10 protein links, or 4 nucleotide neighbors) M23775 Bacteriophage 21 glycoprotein 1 gene, complete cds, and glycoprotein gene, 5' end gi|215451|gb|M23775|PH2GPA [215451] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 3 nucleotide neighbors) M61865 Bacteriophage 21 excisionase (xis), integrase (int) and isocitrate dehydrogenase (icd), complete cds gi|215448|gb|M61865|PH22XISAA [215448] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 protein links, or 9 nucleotide neighbors) S72011 Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds gi|2618967|gb|AF017629|AF017629 [2618967] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors) AF017628 Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds gi|2618964|gb|AF017628|AF017628 [2618964] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors) AF017627 Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds gi|2618961|gb|AF017627|AF017627 [2618961] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors) Bacteriophage 21 isocitrate dehydrogenase (icd) gene, partial cds; and integrase (int) gene, partial cds gi|2618958|gb|AF017626|AF017626 [2618958] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 49 nucleotide neighbors)

WO 00/32825 PCT/IB99/02040

AF017625

# Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds 242 gi|2618955|gb|AF017625|AF017625 [2618955] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors AF017624 Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds gi|2618952|gb|AF017624|AF017624 [2618952] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors) AF017623 Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds gi|2618949|gb|AF017623|AF017623 [2618949] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors) AF017622 Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds gi|2618946|gb|AF017622|AF017622 [2618946] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors) AF017621 Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds gi|2618943|gb|AF017621|AF017621 [2618943] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors) M57455 Bacteriophage 42D (clone pDB17) (from Staphylococcus aureus) staphylokinase gene, complete cds gi|215344|gb|M57455|P42STK [215344] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 protein link, or 9 nucleotide neighbors) Bacteriophage 85 DNA, promoter sequence of unknown gene gi|2058285|emb|Y12633|B85PROM [2058285] (View GenBank report, FASTA report, ASN.1 report, or Graphical view) Bacteriophage P1 DNA sequence around the Op88 operator gi|1359513|emb|X98146|BP1OP88OP [1359513] (View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 nucleotide neighbor) Y07739 Staphylococcus phage Twort holTW, plyTW genes gi|2764979|emb|Y07739|BPTWGHOLG [2764979] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, or 2 protein links) L07580 Bacteriophage phi-11 rinA and rin B genes, required for the activation of Staphylococcal phage phi-11 int expression gil166160|gb|L07580|BPHRINAB [166160] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 2 protein links) Bacteriophage phi-11 integrase (int) and excisionase (xis) genes, complete cds gi|166157|gb|M34832|BPHINTXIS [166157] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 2 nucleotide neighbors)

M20394

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Bacteriophage phi-11 S.aureus attachment site (attP)
     gi|166156|gb|M20394|BPHATTP [166156]
     (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 4 nucleotide neighbors)
  X23128
     Bacteriophage phi-13 integrase gene
     gi|758228|emb|X82312|PHI13INT [758228]
    (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 protein link, or 3 nucleotide neighbors)
  X61719
    S.aureus phi-13 lysogen right chromosome/bacteriophage DNA junction
    gi|46625|emb|X61719|SAP13RJNC [46625]
    (View GenBank report, FASTA report, ASN. 1 report, Graphical view, or 1 MEDLINE link)
 X61718
    S.aureus phi-13 lysogen left chromosomal/bacteriophage DNA junction
   gi|46624|emb|X61718|SAP13LINC [46624]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, or 1 MEDLINE link)
 X61717
   Bacteriophage phi-13 core sequence for attachment
   gi|14799|emb|X61717|BP13ATTP [14799]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE links, or 3 nucleotide neighbors)
 U01875
   Bacteriophage phi-13 putative regulatatory region and integrase (int) gene, partial eds
   gi|437118|gb|U01875|U01875 [437118]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 3 MEDLINE links, or 4 nucleotide neighbors)
X67739
   S.aureus Bacteriophage phi-42 attP gene
   gi|14809|emb|X67739|BPATTPA [14809]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 3 nucleotide neighbors)
U01872
  Bacteriophage phi-42 integrase (int) gene, complete cds
  gi|437115|gb|U01872|U01872 [437115]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 3 MEDLINE links, 2 protein links, or 3 nucleotide neighbors)
X94423
  Staphylococcus aureus bacteriophage phi-42 DNA with ORFs (restriction modification system)
  gi|1771597|emb|X94423|SARMS [1771597]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 protein links, or 1 nucleotide neighbor)
M27965
  Bacteriophage L54a (from S.aureus) int and xis genes, complete cds
  gi|215096|gb|M27965|L54INTXIS [215096]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, MEDLINE 1 link, 2 protein links, or 3 nucleotide neighbors)
U72397
 Bacteriophage 80 alpha holin and amidase genes, complete cds
  gi|1763241|gb|U72397|B8U72397 [1763241]
 (View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 protein links, or 2 nucleotide neighbors)
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AB009866
   Bacteriophage phi PVL proviral DNA, complete sequence
   gi|3341907|dbj|AB009866|AB009866 [3341907]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 63 protein links, or 1 nucleotide neighbor)
 Z47794
   Bacteriophage Cp-1 DNA, complete genome
   gi|2288892|emb|Z47794|BPCP1XX [2288892]
   (View GenBank report, FASTA report, ASN.1 report, Graphical view, 3 MEDLINE links, 28 protein links, 1 nucleotide neighbor, or
   I genome link)
 SEG_CP7RSIT
   Bacteriophage Cp-7 (S.pneumoniae) 5' inverted terminal repeat
   gi|166186|gb||SEG CP7RSIT [166186]
   (View GenBank report, FASTA report, ASN. 1 report, Graphical view, or 1 MEDLINE link)
M11635
   Bacteriophage Cp-7 (S.pneumoniae) DNA, 3' inverted terminal repeat
   gi|166185|gb|M11635|CP7RSIT2 [166185]
   (View GenBank report, FASTA report, ASN.1 report, or Graphical view)
M11636
   Bacteriophage Cp-7 (S.pneumoniae) 5' inverted terminal repeat
   gi|166184|gb|M11636|CP7RSIT1 [166184]
  (View GenBank report, FASTA report, ASN.1 report, or Graphical view)
SEG_CP5RSIT
  Bacteriophage Cp-5 (S.pneumoniae), 5' inverted terminal repeat
  gi|166181|gb||SEG_CP5RSIT [166181]
  (View GenBank report, FASTA report, ASN. 1 report, Graphical view, or 1 MEDLINE link)
M11633
  Bacteriophage Cp-5 (S.pneumoniae) 3' inverted terminal repeat
  gi|166180|gb|M11633|CP5RSIT2 [166180]
  (View GenBank report, FASTA report, ASN.1 report, or Graphical view)
M11634
  Bacteriophage Cp-5 (S.pneumoniae), 5' inverted terminal repeat
  gi|166179|gb|M11634|CP5RSIT1 [166179]
  (View GenBank report, FASTA report, ASN.1 report, or Graphical view)
M34780
  Bacteriophage Cp-9 muramidase (cpl9) gene
  gi|166187|gb|M34780|CP9CPL [166187]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 1 nucleotide neighbor)
M34652
  Bacteriophage HB-3 amidase (hbl) gene, complete cds
  gi|215055|gb|M34652|HB3HBLA [215055]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 protein link)
U64984
  Streptococcus pyogenes phage T12 repressor, excisionase (xis), integrase(int) and erythrogenic toxin A precursor (speA) genes.
  complete cds gij1877426|gb|U40453|SPU40453 [1877426]
  (View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE links, 4 protein links, or 22 nucleotide neighbors)
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WO 00/32825 PCT/IB99/02040

245

Phage CP-T1 (Vibrio cholerae) DNA for packaging signal (pac site) gi|15435|emb|X12375|NCCPPAC [15435]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 protein link)

# AF087814

Vibrio cholerae filamentous bacteriophage fs-2 DNA, complete genome sequence gi|3702207|dbj|AB002632|AB002632 [3702207] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 9 protein links, or 1 genome link)

# D83518

Bacteriophage KVP40 gene for major capsid protein precursor, complete cds gi|3046858|dbj|D83518|D83518 [3046858] (View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, or 1 protein link)

#### AF033322

Bacteriophage PST single-stranded binding protein (gene 32) gene, partial cds, and 5' region gi|2645774|gb|AF033322|AF033322 [2645774] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 protein link, or 17 nucleotide neighbors)

# X94331

Bacteriophage L cro, 24, c2, and c1 genes gi|1469213|emb|X94331|BLCRO24C [1469213] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 4 protein links)

#### U82619

Shigella flexneri bacteriophage V glucosyl transferase (gtr), integrase (int) and excisionase (xis) genes, complete cds gi|2465470|gb|U82619|SFU82619 [2465470] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 8 protein links, or 1 nucleotide neighbor) 246 Table 12

NCBI Entrez Nucleotide QUERY

Key words: bacteriophage and lysis

56 citations found (all selected)

#### AJ011581

Bacteriophage PS119 lysis genes 13, 19, 15, and packaging gene 3, complete cds gil3676084lemblAJ011581lBPS011581 [3676084] (View GenBank report,FASTA report,ASN.1 report,Graphical view,4 protein links, or 1 nucleotide neighbor)

## AJ011580

Bacteriophage PS34 lysis genes 13, 19, 15, antiterminator gene 23, and packaging gene 3, complete cds gil3676078lemblAJ011580lBPS011580 [3676078] (View GenBank report,FASTA report,ASN.1 report,Graphical view,5 protein links, or 2 nucleotide neighbors)

#### AJ011579

Bacteriophage PS3 lysis genes 13, 19, 15, and packaging gene 3 gil3676073|emblAJ011579|BPS011579 [3676073] (View GenBank report,FASTA report,ASN.1 report,Graphical view,4 protein links, or 1 nucleotide neighbor)

# AF034975

Bacteriophage H-19B essential recombination function protein (erf), kil protein (kil), regulatory protein cIII (cIII), protein gp17 (17), N protein (N), cl protein (cl), cro protein (cro), cII protein (cII), O protein (O), P protein (P), ren protein (ren), Roi (roi), Q protein (Q), Shiga-like toxin A (slt-IA) and B (slt-IB) subunits, and putative holin protein (S) genes, complete cds gil2668751|gblAF034975 [2668751]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 20 protein links, or 30 nucleotide neighbors)

# U37314

Bacateriophage lambda Rz1 protein precursor (Rz1) gene, complete cds gil1017780|gblU37314|BLU37314 [1017780] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE links, 1 protein link, or 9 nucleotide neighbors)

# U00005

E. coli hflA locus encoding the hflX, hflK and hflC genes, hfq gene, complete cds; miaA gene, partial cds gil436153lgblU00005IECOHFLA [436153] (View GenBank report,FASTA report,ASN.1 report,Graphical view,4 MEDLINE

links, 5 protein links, or 8 nucleotide neighbors à

#### U32222

Bacteriophage 186, complete sequence gil33372491gblU322221B1U32222 [3337249] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 6 MEDLINE links, 46 protein links, or 5 nucleotide neighbors)

# AF064539

Bacteriophage N15, complete genome gil31926831gblAF064539iAF064539 [3192683] (View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, 60 protein links, 26 nucleotide neighbors, or 1 genome link)

#### AF063097

Bacteriophage P2, complete genome gil3139086|gblAF0630971AF063097 [3139086] (View GenBank report,FASTA report,ASN.1 report,Graphical view,21 MEDLINE links, 42 protein links, 3 nucleotide neighbors, or 1 genome link)

# Z97974

Bacteriophage phiadh lys, hol, intG, rad, and tec genes gil2707950lemblZ97974lBPHIADH [2707950] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE links, 9 protein links, or 1 nucleotide neighbor)

# AF059243

Bacteriophage NL95, complete genome gil30885451gblAF0592431AF059243 [3088545] (View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, 4 protein links, 3 nucleotide neighbors, or 1 genome link)

# AF052431

Bacteriophage M11 A-protein, coat protein, A1-protein, and replicase genes, complete cds
gil2981208igblAF0524311 [2981208]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, 4 protein links, or 8 nucleotide neighbors)

# Y07739

Staphylococcus phage Twort holTW, plyTW genes gil2764979lemblY07739lBPTWGHOLG [2764979] (View GenBank report,FASTA report,ASN.1 report,Graphical view, or 2 protein links)

## X94331

Bacteriophage L cro, 24, c2, and c1 genes gil1469213lemblX94331lBLCRO24C [1469213] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 4 protein links)

#### X78410

Bacteriophage phiadh holin and lysin genes gil793848lembiX78410LGHOLLYS [793848] (View GenBank report,FASTA report,ASN.1 report,Graphical.view,1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)

# X99260

Bacteriophage B103 genomic sequence gil1429229lemblX99260BB103G [1429229] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 17 protein links, or 12 nucleotide neighbors)

# AJ000741

Bacteriophage P1 darA operon gil2462938] emblAJ000741|BPAJ7641 [2462938] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 10 protein links, or 31 nucleotide neighbors)

#### X87420

Bacteriophage ES18 genes 24, c2, cro, c1, 18, and oL and oR operators gil1143407lemblX87420lBPES18GEN [1143407] (View GenBank report,FASTA report,ASN.1 report,Graphical view,5 protein links, or 9 nucleotide neighbors)

### L35561

Bacteriophage phi-105 ORFs 1-3 gil5322181gbiL355611PH5ORFHTR [532218] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 3 protein links)

# D10027

Group II RNA coliphage GA genome gil217784ldbjlD10027lPGAXX [217784] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 3 protein links, 5 nucleotide neighbors, or 1 genome link)

# V01128

Bacteriophage phi-X174 (cs70 mutation) complete genome gil15535iemblV01128iPHIX174 [15535] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 4 MEDLINE links, 11 protein links, or 26 nucleotide neighbors)

#### S81763

coat gene...replicase gene [bacteriophage KU1, host=Escherichia coli, group II RNA phage, Genomic RNA, 3 genes, 120 nt] gil1438766lgblS81763lS81763 [1438766] (View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 MEDLINE link)

#### U38906

Bacteriophage r1t integrase, repressor protein (rro), dUTPase, holin and lysin genes, complete cds gil1353517]gblU38906|BRU38906 [1353517] (View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, 50 protein links, or 3 nucleotide neighbors)

# X91149

Bacteriophage phi-C31 DNA cos region gil1107473|emblX91149|APHIC31C [1107473] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 6 protein links, or 1 nucleotide neighbor)

## V00642

phage MS2 genome gil15081lemblV00642ILEMS2X [15081] (View GenBank report,FASTA report,ASN.1 report,Graphical view,8 MEDLINE links, 4 protein links, or 20 nucleotide neighbors)

# V01146

Genome of bacteriophage T7 gil431187lemblV01146IT7CG [431187] (View GenBank report,FASTA report,ASN.1 report,Graphical view,13 MEDLINE links, 60 protein links, 105 nucleotide neighbors, or 1 genome link)

# X78401

Bacteriophage P22 right operon, orf 48, replication genes 18 and 12, nin region genes, ninG phosphatase, late control gene 23, orf 60, complete cds, late control region, start of lysis gene 13 gil512343lembiX78401POP22NIN [512343] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE links, 13 protein links, or 4 nucleotide neighbors)

# Y00408

Bacteriophage T4 gene t for lysis protein gil15368lemblY00408lMYT4T [15368]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 3 nucleotide neighbors)

## Z26590

Bacteriophage mv4 lysA and lysB genes gil410500lemblZ26590lMV4LYSAB [410500] (View GenBank report,FASTA report,ASN.1 report,Graphical view, or 4 protein links)

#### X07809

Phage phiX174 lysis (E) gene upstream region gil15094lemblX07809lMIPHIXE [15094] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 4 nucleotide neighbors)

# Z34528

Lactococcal bacteriophage c2 lysin gene gil506455lemblZ34528lLBC2LYSIN [506455] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 4 nucleotide neighbors)

# X15031

Bacteriophage fr RNA genome gill 507 llemblX 1503 llLEBFRX [15071] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 4 protein links, 9 nucleotide neighbors, or 1 genome link)

## X80191

Bacteriophage PP7 mRNA for maturation, coat, lysis and replicase proteins gil517237lemblX80191BPP7PR [517237] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 4 protein links, or 1 genome link)

# X85010

Bacteriophage A511 ply511 gene gil853748lemblX85010IBPA511PLY [853748] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 3 protein links, or 1 nucleotide neighbor)

# X85009

Bacteriophage A500 hol500 and ply500 genes gil853744lemblX85009lBPA500PLY [853744] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 3 protein links, or 4 nucleotide neighbors)

### X85008

Bacteriophage A118 hol118 and ply118 genes gil853740/lemblX85008/BPA118PLY [853740] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 3 protein links, or 1 nucleotide neighbor)

### Z35638

Bacteriophage phi-X174 genes for lysis protein and beta-lactamase gil520996lemblZ35638lBPLYSPR [520996]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 516 nucleotide neighbors)

### J02459

Bacteriophage lambda, complete genome gil215104|gblJ02459|LAMCG [215104] (View GenBank report,FASTA report,ASN.1 report,Graphical view,87 MEDLINE links, 67 protein links, 190 nucleotide neighbors, or 1 genome link)

### X87674

Bacteriophage P1 lydA & lydB genes gil974763lemblX87674lBACP1LYD [974763] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 2 nucleotide neighbors)

# X87673

Bacteriophage P1 gene 17
gil974761lemblX87673lBACP117 [974761]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 1 nucleotide neighbor)

### M14784

Bacteriophage T3 strain amNG220B right end, tail fiber protein, lysis protein and DNA packaging proteins, complete cds gil215810gblM14784lPI3RE [215810] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 9 protein links, or 10 nucleotide neighbors)

## M11813

Bacteriophage PZA (from B.subtilis), complete genome gil216046|gblM11813|PZACG [216046] (View GenBank report,FASTA report,ASN.1 report,Graphical view,3 MEDLINE links, 27 protein links, 17 nucleotide neighbors, or 1 genome link)

### M16812

Bacteriophage K3 't' lysis gene, complete cds gil215503lgblM16812lPK3LYST [215503] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 4 nucleotide neighbors)

### J04356

Bacteriophage P22 proteins 15 (complete cds), and 19 (3' end) genes gil215265[gblJ04356|P2215P [215265]

(View GenBank report, FASTA report, ASN. 1 report, Graphical view, 1 MEDLINE link, 3 protein links, or 2 nucleotide neighbors)

#### J04343

Bacteriophage JP34 coat and lysis protein genes, complete cds, and replicase protein gene, 5' end gil215076|gblJ04343lJP3COLY [215076] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 3 protein links, or 2 nucleotide neighbors)

#### J02482

Bacteriophage phi-X174, complete genome gil216019|gbiJ02482iPX1CG [216019]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,23 MEDLINE links, 11 protein links, 26 nucleotide neighbors, or 1 genome link)

#### M99441

Bacteriophage T4 anti-sigma 70 protein (asiA) gene, complete cds and lysis protein, 3' end gil215820lgblM99441lPT4ASIA [215820] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 3 MEDLINE links, 2 protein links, or 2 nucleotide neighbors)

### M65239

Bacteriophage 21 lysis genes S, R, and Rz, complete cds gil215466lgblM65239lPH2LYSGEN [215466] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 3 protein links, or 1 nucleotide neighbor)

### M10637

Phage G4 D/E overlapping gene system, encoding D (morphogenetic) and E (lysis) proteins gil215427|gblM10637|PG4DE [215427] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 12 nucleotide neighbors)

# J02454

Bacteriophage G4, complete genome gil215415[gblJ02454|PG4CG [215415] (View GenBank report,FASTA report,ASN.1 report,Graphical view,6 MEDLINE links, 11 protein links, 20 nucleotide neighbors, or 1 genome link)

### J02580

Bacteriophage PA-2 (E.coli porcine strain isolate) Rz gene, 5'end; ORF2, outer membrane porin protein (lc) and ORF1 genes, complete cds gil215366[gblJ02580]PA2LC [215366] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 4 protein links, or 4 nucleotide neighbors)

#### M14782

Bacillus phage phi-29 head morphogenesis, major head protein, head fiber protein, tail protein, upper collar protein, lower collar protein, pre-neck appendage protein, morphogenesis(13), lysis, morphogenesis(15), encapsidation genes, complete cds gil215323lgblM14782lP29LATE2 [215323] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 11 protein links, or 11 nucleotide neighbors)

#### M10997

Bacteriophage P22 lysis genes 13 and 19, complete cds gil215262lgbiM10997lP221319 [215262] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 3 nucleotide neighbors)

### J02467

Bacteriophage MS2, complete genome gil2152321gbJ02467lMS2CG [215232] (View GenBank report,FASTA report,ASN.1 report,Graphical view,8 MEDLINE links, 4 protein links, 20 nucleotide neighbors, or 1 genome link)

## M14035

Bacteriophage lambda lysis S gene with mutations leading to nonlethality of S in the plasmid pRG1 gil215180lgblM14035ILAMLYS [215180] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 14 nucleotide neighbors)

### U04309

Bacteriophage phi-LC3 putative holin (lysA) gene and putative murein hydrolase (lysB) gene, complete cds gil530796|gblU04309|BPU04309 [530796] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)

Table 13

# NCBI Entrez Nucleotide QUERY

Key word: holin

51 citations found (all selected)

### AF034975

Bacteriophage H-19B essential recombination function protein (erf), kil protein (kil), regulatory protein cIII (cIII), protein gp17 (17), N protein (N), cI protein (cI), cro protein (cro), cII protein (cII), O protein (O), P protein (P), ren protein (ren), Roi (roi), Q protein (Q), Shiga-like toxin A (slt-IA) and B (slt-IB) subunits, and putative holin protein (S) genes, complete cds gil2668751|gblAF034975| [2668751] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 20 protein links, or 30 nucleotide neighbors)

#### U52961

Staphylococcus aureus holin-like protein LrgA (lrgA) and LrgB (lrgB) genes, complete cds gill841516[gblU52961ISAU52961 [1841516] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)

### U28154

Haemophilus somnus cryptic prophage genes, capsid scaffolding protein gene, partial cds, major capsid protein precursor, endonuclease, capsid completion protein, tail synthesis proteins, holin, and lysozyme genes, complete cds gil1765928[gblU28154|HSU28154 [1765928] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 13 protein links)

### AF032122

Streptococcus thermophilus bacteriophage Sfi19 central region of genome gil2935682|gblAF032122| [2935682] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 14 protein links, or 2 nucleotide neighbors)

## AF032121

Streptococcus thermophilus bacteriophage Sfi21 central region of genome gil2935667|gblAF032121|AF032121 [2935667]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 14 protein links, or 2 nucleotide neighbors)

## AF021803

Bacillus subtilis 168 prophage SPbeta N-acetylmuramoyl-L-alanine amidase (blyA), holin-like protein (bhlA), holin-like protein (bhlB), and yolK genes, complete cds; and yolJ gene, partial cds gil2997594|gblAF021803|AF021803 [2997594] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 5 protein links, or 1 nucleotide neighbor)

#### AF057033

Streptococcus thermophilus bacteriophage sfi11 gp502 (orf502), gp284 (orf284), gp129 (orf129), gp193 (orf193), gp119 (orf119), gp348 (orf348), gp53 (orf53), gp113 (orf113), gp104 (orf104), gp114 (orf114), gp128 (orf128), gp168 (orf168), gp117 (orf117), gp105 (orf105), putative minor tail protein (orf1510), putative minor structural protein (orf512), putative minor structural protein (orf570), putative anti-receptor (orf695), putative minor structural protein (orf669), gp149 (orf149), putative holin (orf141), putative holin (orf87), and lysin (orf288) genes, complete cds gil3320432|gblAF057033|AF057033 [3320432] (View GenBank report,FASTA report,ASN.1 report,Graphical view,25 protein links, or 1 nucleotide neighbor)

### U32222

Bacteriophage 186, complete sequence gil3337249[gblU32222|B1U32222 [3337249] (View GenBank report,FASTA report,ASN.1 report,Graphical view,6 MEDLINE links, 46 protein links, or 5 nucleotide neighbors)

### AB009866

Bacteriophage phi PVL proviral DNA, complete sequence gil3341907ldbjlAB009866lAB009866 [3341907] (View GenBank report,FASTA report,ASN.1 report,Graphical view,63 protein links, or 1 nucleotide neighbor)

## AF009630

Bacteriophage bIL170, complete genome gil3282260|gblAF009630|AF009630 [3282260] (View GenBank report,FASTA report,ASN.1 report,Graphical view,63 protein links, 3 nucleotide neighbors, or 1 genome link)

### AF064539

Bacteriophage N15, complete genome

gil3192683|gblAF064539|AF064539 [3192683] (View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, 60 protein links, 26 nucleotide neighbors, or 1 genome link)

### AF063097

Bacteriophage P2, complete genome gil 3139086|gblAF063097|AF063097 [3139086] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 21 MEDLINE links, 42 protein links, 3 nucleotide neighbors, or 1 genome link)

#### Z97974

Bacteriophage phiadh lys, hol, intG, rad, and tec genes gil2707950lemblZ97974lBPHIADH [2707950] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE links, 9 protein links, or 1 nucleotide neighbor)

### X95646

Streptococcus thermophilus bacteriophage Sfi21 DNA; lysogeny module, 8141 bp gil2292747lemblX95646lBSFI21LYS [2292747] (View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, 19 protein links, or 3 nucleotide neighbors)

# SEG\_LLHLYSINO

Bacteriophage LL-H structural protein gene, partial cds; minor structural protein gp61 (g57), unknown protein, unknown protein, structural protein (g20), unknown protein, unknown protein, major capsid protein (g34), main tail protein gp19 (g17), holin (hol), muramidase (mur), unknown protein, unknown protein, unknown protein, unknown protein, unknown protein, unknown protein, unknown protein, unknown protein gene, partial cds; and unknown protein, unknown protein, unknown protein, unknown protein, unknown protein, unknown protein, unknown protein, unknown protein, unknown protein, minor structural protein gp75 (g70), minor structural protein gp89 (g88), minor structural protein gp58 (g71), unknown protein, unknown protein, unknown protein, and unknown protein genes, complete cds gil1004337|gbl|SEG\_LLHLYSINO [1004337]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,4 MEDLINE links, 31 protein links, or 1 nucleotide neighbor)

# M96254

Bacteriophage LL-H holin (hol), muramidase (mur), and unknown protein genes, complete cds gill004336[gblM96254|LLHLYSIN03 [1004336] (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

### Y07740

Staphylococcus phage 187 ply187 and hol187 genes gil2764982lemblY07740lBP187PLYH [2764982] (View GenBank report,FASTA report,ASN.1 report,Graphical view, or 2 protein links)

### U88974

Streptococcus thermophilus bacteriophage 01205 DNA sequence gil2444080|gblU88974| [2444080] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 57 protein links, or 6 nucleotide neighbors)

## Z99117

Bacillus subtilis complete genome (section 14 of 21): from 2599451 to 2812870 gil2634966|emblZ99117|BSUB0014 [2634966] (View GenBank report,FASTA report,ASN.1 report,Graphical view,233 protein links, 51 nucleotide neighbors, or 1 genome link)

### Z99115

Bacillus subtilis complete genome (section 12 of 21): from 2195541 to 2409220 gil2634478|emb|Z99115|BSUB0012 [2634478] (View GenBank report,FASTA report,ASN.1 report,Graphical view,244 protein links, 64 nucleotide neighbors, or 1 genome link)

## Z99110

Bacillus subtilis complete genome (section 7 of 21): from 1194391 to 1411140 gil2633472|emblZ99110|BSUB0007 [2633472] (View GenBank report,FASTA report,ASN.1 report,Graphical view,226 protein links, 31 nucleotide neighbors, or 1 genome link)

## X78410

Bacteriophage phiadh holin and lysin genes gil793848lemblX78410lLGHOLLYS [793848] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)

Z93946

Bacteriophage Dp-1 dph and pal genes and 5 open reading frames gil1934760lemblZ93946lBPDP1ORFS [1934760] (View GenBank report,FASTA report,ASN.1 report,Graphical view, or 6 protein links)

### AF011378

Bacteriophage sk1 complete genome gil2392824|gblAF011378|AF011378 [2392824] (View GenBank report,FASTA report,ASN.1 report,Graphical view,54 protein links, 2 nucleotide neighbors, or 1 genome link)

## Z47794

Bacteriophage Cp-1 DNA, complete genome gil2288892|emb|Z47794|BPCP1XX [2288892]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 3 MEDLINE links, 28 protein links, 1 nucleotide neighbor, or 1 genome link)

## L35561

Bacteriophage phi-105 ORFs 1-3 gil532218|gblL35561|PH5ORFHTR [532218] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 3 protein links)

## D49712

Bacillus licheniformis DNA for ORFs, xpaL2 homologous protein and xpaL1 homologous protein, complete and partial cds gil1514423|dbjlD49712|D49712 [1514423] (View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, or 4 protein links)

# X90511

Lactobacillus bacteriophage phigle DNA for Rorf162, Holin, Lysin, and Rorf175 genes gil1926386|emblX90511|LBPHIHOL [1926386] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 4 protein links, or 1 nucleotide neighbor)

#### X98106

Lactobacillus bacteriophage phigle complete genomic DNA gil1926320lemblX98106lLBPHIG1E [1926320] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE

WO 00/32825 PCT/IB99/02040

link, 50 protein links, or 4 nucleotide neighbors)

#### U72397

Bacteriophage 80 alpha holin and amidase genes, complete cds gil17632411gblU723971B8U72397 [1763241] (View GenBank report,FASTA report,ASN.1 report,Graphical view,2 protein links, or 2 nucleotide neighbors)

### U38906

Bacteriophage r1t integrase, repressor protein (170), dUTPase, holin and lysin genes, complete cds gil1353517igblU38906iBRU38906 [1353517] (View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, 50 protein links, or 3 nucleotide neighbors)

### X91149

Bacteriophage phi-C31 DNA cos region gil1107473|emblX91149|APHIC31C [1107473] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 6 protein links, or 1 nucleotide neighbor)

### U24159

Bacteriophage HP1 strain HP1c1, complete genome gil1046235|gblU24159|BHU24159 [1046235] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 6 MEDLINE links, 41 protein links, 8 nucleotide neighbors, or 1 genome link)

# Z26590

Bacteriophage mv4 lysA and lysB genes gil410500lemblZ26590lMV4LYSAB [410500] (View GenBank report,FASTA report,ASN.1 report,Graphical view, or 4 protein links)

# Z70177

B.subtilis·DNA (28 kb PBSX/skin element region) gil1225934|emblZ70177/BSPBSXSE [1225934] (View GenBank report,FASTA report,ASN.1 report,Graphical view,32 protein links, or 4 nucleotide neighbors)

Z36941

B.subtilis defective prophage PBSX xhlA, xhlB, and xylA genes gil535793lemblZ36941lBSPBSXXHL [535793] (View GenBank report,FASTA report,ASN.1 report,Graphical view,4 protein links, or 5 nucleotide neighbors)

### X89234

L.innocua DNA for phagelysin and holin gene gil1134844lemblX89234lLICPLYHOL [1134844] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 4 nucleotide neighbors)

### X85010

Bacteriophage A511 ply511 gene gil853748|emb|X85010|BPA511PLY [853748] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 3 protein links, or 1 nucleotide neighbor)

### X85009

Bacteriophage A500 hol500 and ply500 genes gil853744lemblX85009lBPA500PLY [853744] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 3 protein links, or 4 nucleotide neighbors)

### X85008

Bacteriophage A118 hol118 and ply118 genes gil853740lemblX85008lBPA118PLY [853740] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 3 protein links, or 1 nucleotide neighbor)

## L34781

Bacteriophage phi 11 holin homologue (ORF3) gene, complete cds and peptidoglycan hydrolase (lytA) gene, partial cds gil511838|gblL34781|BPHHOLIN [511838] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 4 protein links, or 2 nucleotide neighbors)

## U11698

Serratia marcescens SM6 extracellular secretory protein (nucE), putative phage lysozyme (nucD), and transcriptional activator (nucC) genes, complete cds gil509550lgblU11698ISMU11698 [509550] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE

link, 3 protein links, or 1 nucleotide neighbor)

### U31763

Serratia marcescens phage-holin analog protein (regA), putative phage lysozyme (regB), and transcriptional activator (regC) genes, complete cds gil965068lgblU31763lSMU31763 [965068] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 3 protein links, or 1 nucleotide neighbor)

### X87674

Bacteriophage P1 lydA & lydB genes gil974763|emb|X87674|BACP1LYD [974763] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 2 nucleotide neighbors)

# L48605 \

Bacteriophage c2 complete genome gil1146276[gblL48605|C2PVCG [1146276] (View GenBank report,FASTA report,ASN.1 report,Graphical view,3 MEDLINE links, 39 protein links, 3 nucleotide neighbors, or 1 genome link)

### L33769

Bacteriophage bIL67 DNA polymerase subunit (ORF3-5), essential recombination protein (ORF13), lysin (ORF24), minor tail protein (ORF31), terminase subunit (ORF32), holin (ORF37), unknown protein (ORF 1-2,6-12,14-23,25-30,33-36), complete genome gil522252|gblL33769|L67CG [522252] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 37 protein links, 2 nucleotide neighbors, or 1 genome link)

### L31348

Bacteriophage Tuc2009 integrase (int) gene, complete cds; lysin (lys) gene, 3' end gil508612|gblL31348|TU2INT [508612] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE links, 3 protein links, or 3 nucleotide neighbors)

# L31364

Bacteriophage Tuc2009 holin (S) gene, complete cds; lysin (lys) gene, complete cds gil496281lgblL31364lTU2SLYS [496281]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)

### L31366

Bacteriophage Tuc2009 structural protein (mp2) gene, complete cds gil496278|gblL31366|TU2MP2A [496278]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)

## L31365

Bacteriophage Tuc2009 structural protein (mp1) gene, complete cds gil496276[gblL31365]TU2MP1A [496276] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1 protein link)

## U04309

Bacteriophage phi-LC3 putative holin (lysA) gene and putative murein hydrolase (lysB) gene, complete cds gil530796lgblU04309IBPU04309 [530796] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)

### Table 14

NCBI Entrez Nucleotide QUERY Key word: bacteriophage and kil 5 citations found (all selected)

### AF034975

Bacteriophage H-19B essential recombination function protein (erf), kil protein (kil), regulatory protein cIII (cIII), protein gp17 (17), N protein (N), cI protein (cI), cro protein (cro), cII protein (cII), O protein (O), P protein (P), ren protein (ren), Roi (roi), Q protein (Q), Shiga-like toxin A (slt-IA) and B (slt-IB) subunits, and putative holin protein (S) genes, complete cds gil2668751|gblAF034975| [2668751] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 20 protein links, or 30 nucleotide neighbors)

### X15637

Bacteriophage P22 P(L) operon encompassing ral, 17, kil and arf genes gil15646lemblX15637lPOP22PL [15646] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 7 protein links, or 2 nucleotide neighbors)

### J02459

Bacteriophage lambda, complete genome gil215104igblJ02459iLAMCG [215104] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 87 MEDLINE links, 67 protein links, 190 nucleotide neighbors, or 1 genome link)

# M64097

Bacteriophage Mu left end gil215543|gblM64097|PMULEFTEN [215543] (View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE links, 39 protein links, or 15 nucleotide neighbors)

### M18902

Bacteriophage D108 kil gene encoding a replication protein, 3' end; and containing three ORFs, complete cds gil166191lgblM18902lD18KIL [166191] (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 3 nucleotide neighbors)

264

Table 15

U77328	V01282	U11787	U93688	A47599	D21131	U76864	U38428
AF151117	AF121672	U11786	U93687	A47598	D30690	U76863	U66665
AF151218	AF072726	U11785	AJ224764	A47597	D14711	U76862	U66664
AF146368	AF115379	U11784	AF064774	A47596	D90119	U76861	U66663
AF144661	AF034153	U11783	AF064773	A47595	D00730	U76860	X87104
AF132117	AF029244	U11782	Y14370	A47594	D83357	U76859	X87105
Y15477	U67965	U11781	AF065394	A44534	D83356	U76858	X89233
Y09928	U96610	U11780	AF062376	A44533	D83355	U76857	M28521
Y09594	U96609	U11779	AF062375	A44529	D83354	U76855	U54636
AF134905	U73027	U11778	AF062374	A44528	D83353	U76854	U46541
AB019536	U73026	U11777	AF062373	A44527	D12572	U76853	L14017
AJ237696	U73025	U11776	AB007500	A44526	D86727	U76852	U60589
AF106851	AF068904	U11775	Y09924	A44525	D86240	U76851	X48003
AF106850	U60050	U11774	U63529	A39696	D67075	U76850	M37889
AF106849	D10907	U11773	AF033191	AF001783	D67074	U76849	V01281
M26321	D10906	AF053772	Y15856	AF001782	U97062	U76848	X97985
AF060191	AF053140	AF053771	AB000439	L77194	U96620	U76847	X00127
AF060190	AB013298	AF029731	AF041467	AF003593	U96619	Y09929	X03286
AF060189	Y16431	AF027155	Y14051	AF003592	Z84573	Y09570	X62282
AF060188	AF076684	AF024571	U82085	X73889	AB001896	X95848	X01645
AF060187	AF076683	U87144	AF026122	X74219	Y07645	Y09428	X16471
AF060186	Y13225	AF086644	AF026121	Y10419	U92441	S76611	X52734
AF060185	AF094826	AJ223781	AF026120	M63177	U91741	S76213	X13290
AF060184	AJ223480	AF076030	AB009635	E08773	U29454	S75707	X66088
AF036324	AF093548	AF044951	AB006796	E07163	U29478	S75706	Z30588
AF036323	AJ005352	AF044906	U39769	E07162	U77374	S75705	X16457
AF053568	AF051916	AF044905	D00184	E07161	L42945	S76270	X00342
AJ132841	Y09927	AF044904	X56628	E07160	U38429	S72497	V01287
Y13766	AF051917	AF044903	AF033018	E07159	U81980	S72488	X61307
AF101234	S77058	AF044902	AF034076	E07158	X55185	S74031	Y00356
AJ133520	S65052	AF044901	D82063	E07157	V01278	S67449	X06603
AJ133495	AF009671	AF044900	D76414	E07156	U31979	U75367	Z93205
AJ132803	U81973	AF044899	U57060	E07155	X91786	U75368	X64172
AB016487	U77308	AF044898	D89066	E03836	U36912	U31175	X72700
AB016431	U20869	AF044897	U85095	E03835	U36911	X53096	X60827
AB015981	U89396	AF044075	U85097	E03526	U36910	X53951	X64389
AB015195	U94706	AF044074	U85096	E02873	U64885	X53952	X62288
AF107307	U41072	AF044073	D42078	E01690	U76872	X03408	X55798
AF079518	U52961	AF044072	AF015929	E00876	U76871	U50629	X58434
AJ223806	U21636	AF044071	D10369	E00203	U76870	U38656	X06627
Y18018	U65000	AF044070	A48955	D83951	U76869	U58139	X12831
Y17795	U48826	AF044069	A48501	D17366	U76868	A31894	X07371
AJ005647	U20503	AF044068	A48500	D42144	U76867	L42943	X02529
AJ005646	U11789	AF044067	A48499	D42143	U76866	U51474	Y00688
AJ005645	U11788	AF044066	A47600	D10489	U76865	U50077	X04121
X59477	X54338	A12915	U51133	M63176	M10500	L01055	M63917
X59478	X51661	A12913	U51132	L11998	M10499	M83994	M58515
X63598	X05815	A12906	X02588	L05004	AH000934	J03947	L10909
X52593	X15574	A12905	X61716	L42764	M10498	J03479	M15067

WO 00/32825 PCT/IB99/02040

# 265

X76490	Y07536	A12904	X61719	M32103	M10497	M64724	M92376
X81586	X02166	A12903	X61718	U10927	M18264	M14372	M62650
X72014	Z49245	A12902	X67743	AH003057	J01786	M14371	M32312
X72013	X16298	A12901	X67742	M73535	M33833	M14374	M20393
X71437	Z18852	A12900	X67741	M73536	M32470	M15215	M90536
X62992	X68417	A12899	X67740	U20782	M20270	M36694	M21854
X52594	X68425	A12898	X67738	L37598	J03323	M37915	M36771
X14827	X17679	A12897	U02910	L37597	M33479	M12715	L14020
X13404	X63072	A12896	AH003349	L36472	M94061	J04151	M81736
X17301	X02872	A09523	M11118	L25288	M37888	L22566	U11702
X17688	V01277	A04518	M18086	L25893	M76714	L13379	L19300
X03097	X52543	A04517	U19459	K02687	M17123	L13378	L25372
Z16422	A19943	A04512	U35773	L23109	M97169	L13377	L22565
Z33409	A19942	L41499	U26702	L07778	M81346	L13376	M58516
Z33408	A19941	U19770	U21221	M90056	M90693	L13375	U06462
Z33407	A19940	X53818	U36379	J02615	M25257	L13374	L19298
Z33406	A19939	M20129	U06451	M18970	M25256	M17348	M80252
Z33405	A19938	L43098	U35036	K02985	M25255	M17357	L11530
Z33404	A19937	L43082	U20794	M21136	M25254	M17347	
X75439	A19936	X03216	L25426	M10501	M25253	M28364	
X62587	A17958	X70648	M86227	AH000935	M25252	M21319	

- \_\_\_\_\_\_

Table 16

# Phage 44AHJD complete genome sequence. 16668 nucleotides.

1	tccatttct	t tactaaactt	aaaaatgctg	tgcaacaact	taaccaactt	atctaaccta	ttacatattc	
71	atcaaatac	a aaatttatgt	atctattgac	ttttattcaa	aattatgatt	tcaacatata	ataaaattaa	
141	tttacttat	t taaatattct	atgatataat	tagttataaa	atatttggag	gtgtataaat	gacagaattt	
211	gatgaaatc	g taaaaccaga	cgacaaagaa	gaaacttcag	, aatcaactga	agaaaattta	gaatcaactg	
281	aagaaactt	c agaatcaact	gaagaatcaa	ctgaagaato	aactgaagaa	tcaactgaag	ataaaacagt	
351	agaaacaat	c gaagaagaaa	atgaaaacaa	attagaacct	actacaacag	atgaagatag	ttcgaaattt	
421	gaccctgtt	g tattagaaca	acgtattgct	tcattagaac	: aacaagtgac	tacttttta	tcttcacaaa	
491						aacaaagaag		
561						aacatgtatg		
631						gaacagaatt		
701						ttcactttca		
771						aaaaatgaaa		
841		-		_	_	caaaatcttg		
911		•	_			tttattccct		
981	_	-		_		aaagagcaag		
1051						acttatctaa		
1121						cggaattgtg		
1191						gacgcaacta		
1261						aaatgcgtgc		
1331						aaaagaagat		
1401						gaagtacatc		
1471						ttttaacaac		
1541						cattgatttc		
1611						aagttacaaa		
1681 1751						caattccagt		
1821		-				aattaaacca	_	
1891						aaaggtatgt catttaaagc		
1961						ggaagaatta	_	
2031			_		_	caaaggaaat	_	٠
2101						ggaatttta		
2171			_	_	-	tcagctttaa	-	
2241						atgtaaataa		
2311						attaactaaa		
2381						aagctatatg		
2451						atatagaaat		
2521						gcaagcaaaa		
2591				_		atatataacg		
2661		_				caagtaatag		
2731	gcattaactg	aaatgaaacg	ggaatatcaa	aacaaaatta	gtgaattaag	taactattta	ggcattaatt	
2801	cattagccgt	tgataaagaa	agcggtgttt	cagacgaaga	ggcaaaaagt	aatcgtggat	ttaccacatc	
2871	aaacagtaat	atctatttaa	aaggtcgtga	accaattacg	tttttatcaa	agcgttatgg	tttagatatt	
2941	aaaccgtatt	acgatgatga	aacaacgtct	aaaatatcaa	tggtagacac	actttttaaa	gatgaaagca	
3011	gtgatataaa	tggctagata	cacaatgact	ttatacgatt	tcattaaatc	agaattgatt	aaaaaggtt	
3081	tcaatgaatt	tgtaaatgat	aataaattaa	cgttttatga	tgatgaattt	caattcatgc	aaaaaatgct	
3151	gaagttcgac	aaagacgttt	tagctatcgt	taatgaaaaa	gtatttaaag	gtttttcatt	gaaagatgaa	
3221	ttatcagatt	tactttttaa	aaaatcattt	acgattcatt	ttttagatag	agaaatcaac	agacaaacag	
3291				_		tatttaaatg		
3361						atgaagatac		
3431		_	-	-		catgactgca		
3501						acgttacgat		
3571						atcaaaacgc		
3641						aattgataat		
3711						ttacaaattt		
3781	_					cgtccatttt		
3851						ataatactaa		
3921	_	_				tegtatttgg		
3991						ccagagcaag		
4061 4131						agtttaaaaa		
4201					_	gaatgacttt		
4201						ttgataaaga	aaccaaagtt	
4341						taaattaaca		
4411	-	_				ggattagaac		
4481						aagtcgcata		
4551			-		_	gtcactttta	_	
4621						ggtgacggaa		
4691						tgacaattga		
		·						

aacaatgata atcgttggat gcaaggcatt gctgttgatg gtgatgattt atactggtta agtggtaaca 4761 4831 gttcagttaa ttcacatgtt caaatcggta aatattcatt aacaacaggt caaaagattt atgattatcc 4901 atttaagtta tcatatcaag acggtattaa tttcccacgt gataacttta aagagcctga gggtatttgc 4971 atttatacaa atccaaaaac aaaacgtaaa tcgttattac ttgctatgac aaacggcggt ggtggaaaac 5041 gtttccataa tttatatggt ttcttccaac ttggtgagta tgaacacttt gaagcattac gcgcaagagg 5111 ttcacaaaac tataaattaa caaaagacga cggtcgtgca ttatctattc cagaccatat cgacgattta 5181 aatgacttaa cgcaagctgg tttttattat attgacgggg gtactgcaga aaaacttaag aatatgccaa 5251 tgaatggtag caagcgtata attgacgctg gttgtttcat taatgtatac cctacaacac aaacattagg 5321 tacggttcaa gaattaacac gtttctcaac aggtcgtaaa atggttaaaa tggtgcgtgg tatgacttta 5391 gacgtattta cgttaaaatg ggattatgga ttatggacaa caatcaaaac tgacgcacca tatcaagaat 5461 attrggaage aagteaatae aataactgga ttgettatgt aacaacaget ggtgagtatt acattacagg taaccaaatg gaattattta gagacgcgcc agaagaaatt aaaaaagtgg gtgcatggtt acgtgtgtca 5531 agtggtaacg 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gcagtttttg aagatggttc gtttttagtt gcaaactata atgtaccacc atatgttgca ccatcacgtg 10081 10151 tggtattgta tacactcatt aatggegtac caaataatgc tggtgataat attgtattct ttagtggtat 10221 tgcttaatta actatgctat aatgaacaca tgctagtaat gctagtaaat aaaatacaaa acataatcaa ttttcgtaca catttttcat gttatctcaa aaagaaaagg agactgttat tttaacagtt gcctttttt 10291 10361 atttcatcat gttcacgttt taatatatgc aaatcagatt tgttatgtac tgaacgttca actggaaata 10431 aqteqttaaq tqaaaatqaa ccqatqteac tttcaatata aagaatatca tcaaattqac tatqqteqaa 10501 atttteteta gegtetttta atataaatte aegttteata ttaagtteat eagtaaaata tteateatat agattaccac atacaatttc agttttagac qqatatatcq atattqtacc ttqctcatta taqatacttt 10571 10641 tattgttttc aataatggca ccgtcaaaga attgttcacg tacaaaggtt tcaaaatcga cgcttgtatc 10711 aaaggcgttt ttcggtatac cagcagaagc aattttaatc tttccattca cttcatatgc atatttctta 10781 tgattcagta caaacatctt atctatctgt tcgttttcaa tatcccattt acctaaggct atcgggtcga ataaactggg gttcaataag ggtttaacaa cggatttcat atacaaacta tcagtatcgc aataaataaa 10851 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15401	cagtgacgat	aacaccttgt	tcaccgaatt	ttgattcttt	gtttgtgaat	aatgctctaa	cgatatactc
15471	ttttttcata	ccgtatttt	ctactaattc	tgatagtttg	ataaattctc	tttcttttc	ctcaaattca
15541	aatctcgcta	atgtgttttg	gtgtcttgat	aaaatatctt	ttacgtttgt	cattttattt	ctcctcttat
15611	ttaaattatt	tgctttctgc	aattgcgatt	tgtagtaaat	cattgtaata	aacttgaatt	gttttcgttg
15681	tgcgtgtagt	ggacaatagt	ttacatgtgt	ctggtaataa	ttcttttgct	tgtgttttgg	ttaaatgata
15751	ctcgtgaagt	ggtaaaaatt	cctcaatgta	ttcattatca	tcatctaagt	aatgaagtat	ataacctttg
15821	acacgtaagg	taacaatgtc	gtcaactttc	attattatat	cactcctttc	taaaaaacgt	aaacgttata
15891	cgtttcataa	aatcctttat	gcatattcca	ttgttctatt	gggtcatcac	cagcaatata	agacaatatt
15961	gattetggtt	tagtttcgtt	gtttagttca	tcatttaaga	attgaacaac	agaactatta	tagtttaata
16031	atagttgttg	gcaagccgat	aataagttaa	ttgcattgtc	aaatgtataa	gctggattcc	attgaatcag
16101	tttattgaat	agttgcaaca	tttcagtata	ggcttgtcct	ttttcttctg	gtgcattatc	aacattaacc
16171	attattatca	cttcctaata	aagttgaaat	tacgcgtaaa	acagaattat	gatttaaatc	ttcaatttca
16241	tcaatqtcaa	catcataaaa	tgaaatttca	ttttctgttc	tatcaaataa	cgctatacat	aaacttccat
16311	tettaaaacg	aaaaacatgc	ttcaactcaa	tgttttttgt	ttcattttcc	atttttgtta	ctccttgttt
16381					caatagtttt		
16451					ttcaaatcat		
16521	agatacataa	attttgtatt	tgatgaatat	gtaataggtt	agataagttg	gttäagttgt	tgcacagtat
16591					tttgatttgt		
16661	ggtggggt			_		_	

- \_\_\_\_\_\_\_\_

Table 17

# Phage 44AHJD ORFs list

nb	Name	Frame	Position	Size (a.a.)	Key words
1	44AHJDORF001	-1	1034212627	761	DNA polymerase;
2	44AHJDORF002	3	37895732	647	Techoic acid; Staph;
3	44AHJDORF003	2	66268389	587	Tail:
4	44AHJDORF004	1	876410227	487	Serine protease motif;
5	44AHJDORF005	-1	1264313890	415	
6	44AHJDORF006	2	8032029	408	
7	44AHJDORF007	1	20443027	327	Upper collar;
8	44AHJDORF008	2	30203775	251	Lower collar;
9	44AHJDORF009	2	57446496	250	Amidase; Staph;
10	44AHJDORF010	-2	1393814420	160	
11	44AHJDORF012	3	83918813	140	Holin;
12	44AHJDORF013	-2	1458614996	136	
13	44AHJDORF113	1 1	199600	133	
14	44AHJDORF011	-2	1522515593	122	
15 16	44AHJDORF114 44AHJDORF014	-2	1587016172	100 92	<del> </del>
17	44AHJDORF015	1 1	62436521 1540315645	80	
18	44AHJDORF016	-1	1561615852	78	
19	44AHJDORF017	-2	1053610757	73	
20	44AHJDORF018	-1	8861098	70	<del>-</del>
21	44AHJDORF019	-2	96309836	68	
22	44AHJDORF121	-1	1616516362	65	
23	44AHJDORF020	2	1386514053	62	
24	44AHJDORF123	2	614796	60	
25	44AHJDORF021	-2	56345816	60	
26	44AHJDORF023	-2	63156494	59	
27	44AHJDORF024	1	1427514451	58	
28	44AHJDORF025	-3	1499915175	58	
29	44AHJDORF026	-3	1442614593	55	
30	44AHJDORF027	1	1291613080	54	
31	44AHJDORF029	-1	1501915183	54	
32 33	44AHJDORF028	3	90719235	54	
34	44AHJDORF030 44AHJDORF031	2	1448714648 1103911191	50	
35	44AHJDORF135	3	693842	49	
36	44AHJDORF033	1 -1	36463795	49	· · · · · · · · · · · · · · · · · · ·
37	44AHJDORF032	-2	93069455	49	
38	44AHJDORF034	-3	1400014146	48	
39	44AHJDORF035	-3	1381113957	48	
40	44AHJDORF036	-3	1001910165	48	
41	44AHJDORF022	-3	84688611	47	
42	44AHJDORF037	1	1478814931	47	
43	44AHJDORF038	-2	35283671	47	
44	44AHJDORF039	3	17431883	46	
45	44AHJDORF040	2	97409877	45	
46 47	44AHJDORF041	2	1583615973	45	
48	44AHJDORF042 44AHJDORF043	-1 -1	50145151 44024539	45 45	
49	44AHJDORF044	-2	1278312917	44	
50	44AHJDORF149	-2	639770	43	
51	44AHJDORF046	1	48915019	42	
52	44AHJDORF047	1 1	1191112039	42	
53	44AHJDORF045	2	1065510783	42	
54	44AHJDORF048	-3	1521215340	42	
55	44AHJDORF049	3	57845909	41	
56	44AHJDORF050	3	1315813283	41	
57	44AHJDORF051	-2	1094411066	40	
58	44AHJDORF052	-3	1421614338	40	-
59	44AHJDORF053	3	33483467	39	
50	44AHJDORF054	3	75517670	39	
51	44AHJDORF055	3	1570515821	38	
52	44AHJDORF056	1	55125625	37	
3	44AHJDORF057	2	1012110231	36	
54	44AHJDORF058	3	1076710877	36	

WO 00/32825

65	44AHJDORF164	-1	592702	36	
		-			
66	44AHJDORF059	-2	82508360	36	
67	44AHJDORF060	-2	61476257	36	
68	44AHJDORF061	2	1555115658	35	
69_	44AHJDORF062	1	42854389	34	
70	44AHJDORF063	-3	93839487	34	
71	44AHJDORF065	1 1	50295130	33	
72	44AHJDORF064	2	26092710	33	
73	44AHJDORF066	-2	1038010481	33	

WO 00/32825 PCT/IB99/02040

272

# Table 18

# Predicted amino acid sequences

```
44AHJDORF001
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12627
     MGLLBCMQYHKHERRMILYWDIETLA
     12543
      KVNGRKKPTKYKNVTYSVAIGWFNGYEI
29
     qatgttgaagtatttccgagtttcgaatctttttatgacgcattttatacgtatgtgaaaagacgtgatacaatcacaaaatca
12459
      DVEV PPS FES FYD A FYTYV KRRDTITKS
     12375
      K T D I I M I A H N C N K Y D N H F L L K D T M R Y F
85
     aatattacacgcgaaaatatatatttaaaatctgcagaagaaaatgaacacacattaaaaatgaaagaggctactattttagcc
12291
      NITRENIYLKSAEENEHTLKMKEATILA
113
     12207
      K N Q N V I L E K R V K S S I N L D L T M F L N G F K F
141
     aatattattgataactttatgaaaaccaatacatcaattgcaacattaggtaagaaattacttgatggtggttatttaacagaa
12123
      NIIDNFMKTNTSIATLGKKLLDGGYLTE
169
      tcacaacttaaaacagattttaattatacgatttttgataaagataatgatatgatagtgaagcctatgactatgctgtg
12039
      SQLKTDFNYTIFDKDNDMNDSEAYDYA
     11955
225
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11871
      SDIFPNFDYNKLTFSLNIMESYLNNEM
253
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       a a a traac g tratag at g a tratag ca a tratag ca a a a a tratag ca a a a tratag ca a a a tratag ca a a tratag ca a a tratag ca a tratag ca a a tratag ca a a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tratag ca a tr
        K L T F Y D D E F Q F M Q K M L K F D K D V L A I V N E
3188
       K V F K G F S L K D E L S D L L F K K S F T I H F L D R
3272
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3356
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113
3440
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3524
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169
        E V N I D V D N T T L R F A D N N T I D N G K T V N-LK'S
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S N E S N Q N A K R N Q N Q K G N A K G T Q F T K Q Y L
3608
197
3692
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3775
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225
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44AHJDORF009

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5828
       C M D L S V A Y V Y Y I T D G K V R M W G N A K D A I N
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5912
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57
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6416
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14252
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14084
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113
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141
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       15509
       K L S E L V E K Y G M K K E Y I V R A L F T N K E S K F
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15257
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8475
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8559
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57
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8643
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8727
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       taa 8813
8811
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14912
       29
14828
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14744
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44AHJDORF113
199
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       M T E F D E I V K P D D K E E T S E S T E E N L E S T E ...
283
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367
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57
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451
       85
       S L E Q Q V T T F L S S Q M Q Q P Q Q V Q Q T Q S D V T
535
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16088
      N P A Y T F D N A I N L L S A C Q Q L L L N Y N S S V V
29
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15920
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      M K M V H L H V V F Y Q Y L H V S V V Q N Y Q N L M A I
6327
      G S N Q T V I H H I T K F V Y Q M V T Y G L V I T G K A
29
6411
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6495
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85
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15487
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15768
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9668
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     M Y E G N N M R S M M G T S Y E D S R L N K R T E L N E
698
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782
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57
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14443 cataaataa 14451
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15019
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9235
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44AHJDORF033
3795
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29
44AHJDORF035
13957 atgcaacatttgacgaataaatttaacactgtaaacgacatcataaactattacaaggagcaaaaacatggtaaaacaaaatcg
      MQHLTNKFNTVNDIINYYKEQKHGKTKS
13873 tttagacatggtaagagattatcaaaatgctgtcaatcatgtcagaaaaaaatcccagataa 13811
      FRHGKRLSKCCQSCQKKNPR*
44AHJDORF036
{\tt 10165} \quad {\tt gtgtatacaataccacacgtgatggtgcaacatatggtggtacattatagtttgcaactaaaaacgaaccatcttcaaaaactg}
      V Y T I P H V M V Q H M V V H Y S L Q L K T N H L Q K L
10081 ctacaacaacacctgtgtgaccaataccatatgcagttgcttgtaagtatggtggtttactag 10019
29
      L Q Q H L C D Q Y H M Q L L V S M V
44AHJDORF037
1 M S I S N V N N S F S I S K S S Y C L S N S I Y T S P I 14872 tttattttattatatatatataaaatttttcaaatttttcaaatttttcaaatttttcatataa 14931
29
      FIFTIHFLLDEVNFSNLSL
44AHJDORF038
      gtgtacctttttgcattacctttttgattttgattacgttttgcgttttgattactttcgttactcgatttattcacagttttac
V Y L L H Y L F D F D Y V L R F D Y F R Y S I Y S Q F Y
cgttatcaatcgtattattatcagcgaatcgtaacgttgtattatcaacatcaatgttaa 3528
R Y Q S Y Y Y Q R I V T L Y Y Q H Q C *
3671
3587
29
44AHJDÖRF039
      \tt gtgctgtatttacttatgatgtatctaaacttaaagagtttactggcaacgttgaagaaattaaaccaaaatcagatttatatg
1743
      V L Y L L M M Y L N L K S L L A T L K K L N Q N Q I Y M
      cgtttattttggatattaattcaattaaatataaacgttacacaaaaggtatgttaa 1883
      RLFWILIQLNINVTQK
44AHJDORF040
      gtggtaactggacatatgcacagttaccagaaaaatataaaaaagcaattggtgtacctttattcaaaaaagaatacttataca
      V T G H M H S Y Q K N I K K Q L V Y L Y S K K N T Y T
9824
      aaccaggtaacatatttcctcaaacgggtaatgcaggacaatgtacagaattaa 9877
      NQVTYFLKRVMQDNVQN
29
44AHJDORF041
15836 atgregreaactttcattattatcactcctttctaaaaaacgtaaacgttatacgtttcataaaatcctttatgcatattcc
      M S S T F I I I S L L S K K R K R Y T F H K I L Y A Y S
15920
      attgttctattgggtcatcaccagcaatataagacaatattgattctggtttag 15973
29
      I V L L G H H Q Q Y K T I L I L V *
44AHJDORF042
5151
      {\tt atgcacqaccqtcgtcttttgttaatttatagttttgtgaacctctttgcgcgtaatgcttcaaagtgttcatactcaccaagtt}
      M H D R R L L L I Y S F V N L L R V M L Q S V H T H Q V
5067
      ggaagaaaccatataaattatggaaacgttttccaccaccgccgtttgtcatag 5014
29
      GRNHINYGNVFHHRRLS*
44AHJDORF043
4539
      atgcgacttgtaacagttttgcaacaccatcgtgatgtaaccagattttcatttcaccattggattgacgttctaatccgattg
      M R L V T V L Q H H R D V T R F S F H H W I D V L I R L ttgtaccatgaccaccctgtacaatacgcatgcttgaaattaagtcaccactag 4402
4455
      LYHDHPVQYACLKLSHH
29
44AHJDORF044
      \verb|atgttacctatttacgtgatgatatgttttataaagaaaacatggaacgttattactacaatccaagcaatttacattttgaca| |
12917
      MLPIYVMICPIKKTWNVITTIQAIYILT
      M L T L K I T W L I M I D I Y I *
44AHJDORF149
      {\tt atgattgttttgaaagtgaatgaatttgtacaccataactatcttcacttttatttgtatcaattgacatgttttcatttaatt}
      MIVLKVNEFVHHNYLHFYLYQLTCFHLI
686
      ctgttcgtttatttaatcttgaatcttcatatgatgtacccatcatag 639
29
      LFVYLILNLHMMYPS
44AHJDORF046
      {\tt atgattatccatttaagttatcatatcaagacggtattaatttcccacgtgataactttaaagagcctgagggtatttgcattt}
4891
      M I I H L S Y H I K T V L I S H V I T L K S L R V-F A F
4975
      atacaaatccaaaaacaaaacgtaaatcgttattacttgctatga 5019
      IQIQKQNVNRYYLL
29
44AHJDORF047
```

WO 00/32825 PCT/IB99/02040

```
11995 atctttatcaaaaatcgtataattaaaatctgttttaagttgtga 12039
      I F I K N R I I K I C F K L
44AHJDORF045
10655 atggcaccgtcaaagaattgttcacgtacaaaggtttcaaaatcgacgcttgtatcaaaggcgtttttcggtataccagcagaa
1 MAPSKNCSRTKVSKSTLVSKAFFGIPAE
10739 gcaattttaatctttccattcacttcatatgcatatttcttatga 10783
       A I L I F P F T S Y A Y F L
29
44AHJDORF048
{\tt 15340} \quad {\tt atgaggacgttgttgacattatcaatgctggagaagttcaattcacaatttatgaatatgaaaacaaaaaggtcaaaaaggtt}
1 MRTLLTLSMLEKFNSQFMNMKTKKVKK
15256 actcaatcaattttggtcaagtatcattttaatacaatttcatag 15212
      TQSILVKYHFNTIS .
44AHJDORF049
      {\tt atgagggggeaggtgttgactttgatggtgcatatggatttcaatgtatggacttatcagttgcttatgtgtattacattactg}
5784
       M R G Q V L T L M V H M D F N V W T Y Q L L M C I T L L
       acggtaaagttcgcatgtggggtaatgctaaagacgcgataa 5909
5868
       TVKFACGVMLKTR
29
44AHJDORF050
     gtgtgttacgtttttcattcacgtaatcgtttcgtcgcatttctaaaaaaatgtttttgtaaagtcttgatgtattcattttat
13158
       V C Y V F H S R N R F V A F L K K C F C K V L M Y S F Y
13242 gcttttgtaataaattgtatatatttaaattggataatatag 13283
      A F V I N C I Y L N W I I *
29
44AHJDORF051
     atgataacaatgaactatacaatatcattaacggttacaaaaacactgaacgtaatatattattctctacatttgtcacatcac
11066
       MITMNYTISLTVTKTLNVIYYSLHLSHH
10982 gttcattgtataacttattggttcctttccaatacttaa 10944
       VHCITYWFLSNT
44AHJDORF052
14338 atgattttagtaatgttaattttaaatttgatgataaagatttacaagaggcgtacattgacacatggaaacatttttgcacatc
1 M I L V M L I L N L M I K I Y K R R T L T H G N I L H I
14254 tgccctattttcctaaagaaagaaacgtatcatatgtaa 14216
       CPIFLKKETYHM
29
44AHJDORF053
      at \verb|gtggtttattcatcaagtgaagttgaaaaatacttacaatcacaaggcttcacagaacacaatgaagatacaacaagtaaca
3348
      M W F I H Q V K L K N T Y N H K A S Q N T M K I Q Q V T ctgatgaaacatcgaatcaaaatgctacatctttag 3467
3432
29
       LMKHRIKMLHL
44 NH.TDORROS4
      atgactggaatggaaatacgatgttactcgacgctggtaagatttcacaaaaaactggtgttaagttacgtacaaaatcaatta
мт с м е і к с у s т L v к f н к к L v L s у v Q N Q L
7551
       ttggttatcataatgaagttcgagtatatccagtag 7670
7635
       LVIIMKFEYIQ
44AHJDORF055
15705 atgtgtctggtaataattcttttgcttgtgttttggttaaatgatactcgtgaagtggtaaaaattcctcaatgtattcattat
       M C L V I I L L V F W L N D T R B V V K I P Q C I H Y
15789 catcatctaagtaatgaagtatataacctttga 15821
       HHLSNEVYNL
44AHJDORF056
      gtgagtattacattacaggtaaccaaatggaattatttagagacgcgccagaagaaattaaaaaagtgggtgcatggttacgtg
      V S I T L Q V T K W N Y L E T R Q K K L K K W V H G Y V tgtcaagtggtaacgcagtcggtgaagtaa 5625
       CQVVTQSVK *
44AHJDORF057
{\tt 10121} \quad {\tt atgtaccaccatatgttgcaccatcacgtgtggtattgtatacactcattaatggcgtaccaaataatgctggtgataatattg}
     MYHHMLHHHVWYCIHSLMAYQIMLVII L
tattctttagtggtattgcttaattaa 10231
10205
29
       Y S L V V L L N *
44AHJDORF058
     10767
1
10851 ataaactggggttcaataagggtttaa 10877
       INWGSIRV *
29
44AHJDORF164
       at {\tt gttttcatttaattctgttcgtttatttaatcttgaatcttcatatgatgtacccatcatagaacgcatgttgtttccctca}
702
       MFSFNSVRLFNLESSYDVPIIERMLFPS
1
       tacatgtttaaattcctcctaatctaa 592
618
       YMFKFLLI
29
44AHJDORF059
       atggattttgtaacattggattacctgaaccgtcattatgccaaaatcttacaccagattctaaaattgcttttaattgttcca
8360
       M D F V T L D Y L N R H Y A K I L H Q I L K L L & I V P
       ttaacatggggtcgatgtcacgtatag 8250
8276
       LTWGRCHV
44AHJDORF060
       atgtaccattttcatttctataatatgtgccgtattggtttcgtttccattttccaaatgtatttacttttgatgtttctaatg
```

```
6173 ctttgctattactacctgaaaatttag 6147
      rcaaark:
44AHJDORF061
{\tt 15551} \quad {\tt atgtgttttggtgtcttgataaaatatcttttacgtttgtcattttatttctccctcttatttaaattatttgctttctgcaatt}
      M C F G V L I K Y L L R L S F Y P S S Y L N Y L L S A I
15635
     gcgatttgtagtaaatcattgtaa 15658
29
      A I C S K S L *
44AHJDORF062
      4285
1
      aacctgaaggtttttggataa 4389
4369
29
      N L K V F G *
44AHJDORF063
9487
     atgogtottgtattttttttaataattottgcatggottgttttgctaaagcgagtagtgaactaccactgtcaccactactac
1
      M R L V F F L I I L A W L V L L K R V V N Y H C H H Y Y
9403
      cactgtcagacgaatcactag 9383
29
      H C Q T N H *
44AHJDORF065
      gtggtggaaaacgtttccataatttatatggtttcttccaacttggtgagtatgaacactttgagcattacgcgcaagaggtt V V E N V S I I Y M V S S N L V S M N T L K H Y A Q E V cacaaaactataaattaa 5130
5029
1
5113
      H K T I N *
29
44AHJDORF064
     atgacgagtcaatcaatcaacttgtgtccgaaatatataacggtgcaccatttgttaaaatgtcacctatgtttaatgcagatg
2609
      M T S Q S I N L C P K Y I T V H H L L K C H L C L M Q M acgatatcattgatttaa 2710
29
      TISLI
44AHJDORF066
10481 \quad atgatattctttatattgaaagtgacatcggttcattttcacttaacgacttatttccaqttgaacqttcagtacataacaaat
     MIFFILKVTSVHFHLTTYFQLNVQYITN
10397 ctgatttgcatatattaa 10380
     LICIY*
```

#### Table 19

# Sequence similarities between ORFs 44AHJD and public databases

```
Phage: 44AHJD
Database: nr
Query= sid|110871|lan|44AHJDORF001 Phage 44AHJD ORF|10342-12627|-1
            (761 letters)
gi|118848|sp|P19894|DPOL_BPM2 DNA POLYMERASE >gi|76896|pir||JQ0...
                                                                                          55 le-06
gi|1072656|pir||S51275 DNA polymerase - phage CP-1 >gi|836593|e...
                                                                                               6e-06
gi|1429230|emb|CAA67649| (X99260) DNA polymerase [Bacteriophage...
gi|1572479|emb|CAA65712| (X96987) DNA polymerase [Bacteriophage...
gi|118851|sp|P06950|DPOL_BPPZA DNA POLYMERASE (EARLY PROTEIN GP...
                                                                                          49 1e-04
                                                                                          46
                                                                                               0.001
                                                                                               0.002
gi|2435429 (AF012250) unassigned reading frame (possible DNA po...
gi|1084487|pir||S41618 DNA polymerase - slime mold (Physarum po...
                                                                                               0.002
                                                                                               0.002
gi|4877819|gb|AAD31446.1| (AP133505) DNA polymerase [Neurospora...
                                                                                          44
gi|461962|sp|P33537|DPOM NEUCR PROBABLE DNA POLYMERASE >gi|2833...
gi|2499511|sp|012471|6P22 YEAST 6-PHOSPHOFRUCTO-2-KINASE 2 (PHO...
gi|2258375|gb|AAD11909.1| (AF007261) transcription initiation f...
                                                                                          44 0.004
                                                                                          41 0.041
                                                                                          40 0.070
gi 15734 emb CAA37450 (X53370) DNA polymerase (AA 1-575) [Bact...
                                                                                          39 0.092
Query= sid|110872|lan|44AHJDORF002 Phage 44AHJD ORF|3789-5732|3
            (647 letters)
gi|135273|sp|P27622|TAGC_BACSU TEICHOIC ACID BIOSYNTHESIS PROTE...
                                                                                         112 7e-24
gi 142847 (M64050) DNase inhibitor (Bacillus subtilis)
                                                                                          52 1e-05
gi 4038407 (AF103943) factor C protein precursor (Streptomyces ...
                                                                                          39 0.10
Query= sid|110873|lan|44AHJDORF003 Phage 44AHJD ORF|6626-8389|2
            (587 letters)
gi|138123|sp|P04331|VG9_BPPH2 TAIL PROTEIN (LATE PROTEIN GP9) >...
                                                                                          92 8e-18
gi|138124|sp|P07534|VG9_BPPZA TAIL PROTEIN (LATE PROTEIN GP9) >...
                                                                                          82 1e-14
gi|1429238|emb|CAA67657| (X99260) tail protein [Bacteriophage B...
gi|215339 (M12456) pp tail protein [Bacteriophage phi-29] >gi|1181968|emb|CAA87738.1| (Z47794) tail protein [Bacteriophage...
gi|1181970|emb|CAA87740.1| (Z47794) tail protein [Bacteriophage...
                                                                                          78 2e-13
                                                                                          71 2e-11
                                                                                          54 3e-06
                                                                                          42 0.010
Query= sid|110875|lan|44AHJDORF005 Phage 44AHJD ORF|12643-13890|-1
            (415 letters)
gi|3845203 (AE001399) GAF domain protein (cyclic nt signal tran...
                                                                                          52 6e-06
gi|3758843|emb|CAB11128.1| (Z98551) predicted using hexExon; MA...
gi|3845297 (AE001421) hypothetical protein [Plasmodium falciparum]
                                                                                               5e-05
                                                                                          48 le-04
gi 4493936 emb CAB38972.1 (AL034556) predicted using hexExon; ...
                                                                                           47
                                                                                               2e-04
gi 3845165 (AE001390) hypothetical protein (Plasmodium falciparum)
                                                                                          46 6e-04
Query= sid | 110877 | lan | 44AHJDORF007 | Phage 44AHJD ORF | 2044-3027 | 1
            (327 letters)
gi|1181960|emb|CAA87731.1| (Z47794) connector protein [Bacterio...
                                                                                          46 Se-04
gi 1429239 emb CAA67658 (X99260) upper collar protein [Bacteri...
                                                                                          45 8e-04
gi | 137915 | sp | P07535 | VG10 BPPZA UPPER COLLAR PROTEIN (CONNECTOR ...
                                                                                          44 0.002
                                                                                          41 0.009
gi 137914 BP P04332 VG10 BPPH2 UPPER COLLAR PROTEIN (CONNECTOR ...
Query= sid|110878|lan|44AHJDORF008 Phage 44AHJD ORF|3020-3775|2
            (251 letters)
gi|4982468|gb|AAD30963.2| (AF118151) SNF1/AMP-activated kinase ...
                                                                                          52 3e-06
gi|1730077 sp|P18160|KYK1_DICDI NON-RECEPTOR TYROSINE KINASE SP...
                                                                                          46 2e-04
gi|3758855|emb|CAB11140.1| (Z98551) predicted using hexexon; MA... gi|585795|sp|P21538|REB1_YEAST DNA-BINDING PROTEIN REB1 (QBP) >...
                                                                                          46
                                                                                               2e-04
                                                                                          46
                                                                                               3e-04
                                                                                                        <u>____</u>
gi|172372 (M58728) DNA-binding protein [Saccharomyces cerevisiae]
                                                                                          46 3e-04
gi|2952545 (AF051898) coronin binding protein [Dictyostelium di...
gi|535260|emb|CAA82996| (Z30339) STARP antigen [Plasmodium reic...
                                                                                          45 6e-04 _
                                                                                          45 7e-04
gi|1429240|emb|CAA67659| (X99260) lower collar protein [Bacteri...
                                                                                          44 0.001
```

# Query= sid|110879|lan|44AHJDORF009 Phage 44AHJD ORF|5744-6496|2 (250 letters)

gi 2764981 emb CAA69021.1  (Y07739) N-acetylmuramoyl-L-a	lanine 18	0 le-44
gi 113675 sp P24556 ALYS_STAAU AUTOLYSIN (N-ACETYLMURAMO	YL-L-AL 11	8 6e-26
gi 1763243 (U72397) amidase [bacteriophage 80 alpha]	11	8 6e-26
gi 4574237 gb AAD23962.1 AF106851 1 (AF106851) LytN (Sta	phyloco 8	4 9e-16
gi 3767593 dbj BAA33856.1 (AB015195) Lyth (Staphylococci	us aureus) 8	4 9e-16
gi 2764983 emb CAA69022.1  (Y07740) cell wall hydrolase	Ply187 7	7 2e-13
gi 3287732 sp 005156 ALE1_STACP GLYCYL-GLYCINE ENDOPEPTI	DASE AL 7	3 2e-12
gi 79926 pir  A25881 lysostaphin precursor - Staphylococ	cus sim 6	9 3e-11
gi   126496   sp   P10548   LSTP_STAST LYSOSTAPHIN PRECURSOR (GL	YCYL-GL 6	9 3e-11
gi 3287967 sp P10547 LSTP_STASI LYSOSTAPHIN PRECURSOR (G		9 3e-11
gi 3341932 dbj BAA31898.1  (AB009866) amidase (peptidogly	ycan hy 6	8 6e-11
Query= sid 110882 lan 44AHJDORF012 Phage 44AHJD ORF 8391 (140 letters)	-8813 3	
qi 140528 sp P24811 YQXH BACSU HYPOTHETICAL 15.7 KD PROT	EIN IN 80	0 6e-15
qi 4126631 db  BAA36651.1  (AB016282) ORF45 (bacteriopha		
qi 141088 sp P26835 YNGD CLOPE HYPOTHETICAL 14.9 KD PROTI		
qi 2293160 (AF008220) YtkC [Bacillus subtilis] >qi 263554		
qi 1181973 emb CAA87743.1  (Z47794) holin protein [Bacter		
atimoty, a completely to the form the property the property the property that the pr	,10paag J.	. 3.3

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#### Table 20

Homolgies between phage 44 AHJD ORFs and proteins in public databases

```
Query= pt | 110871 44AHJDORF001 Phage 44AHJD ORF | 10342-12627 | -1 1
          (761 letters)
>gi|118848|sp|P19894|DPOL_BPM2 DNA POLYMERASE >gi|76896|pir||JQ0161
             DNA-directed DNA polymerase (EC 2.7.7.7) - phage M2
             >gi|215509 (M33144) DNA polymerase [Bacteriophage M2]
             Length = 572
 Score = 55.4 bits (131), Expect = 1e-06
 Identities = 96/426 (22%), Positives = 159/426 (36%), Gaps = 88/426 (20%)
Query: 229 KLTPEQLTYIHNDVIILGMCHIHYSDIFPNFDYNKLTFSLNIMESYLNNEMTR-----FQ 283
++TPE+ YI ND+ I+ DI +++T + ++ + T+ F
Sbjct: 154 EITPEEYEYIKNDIEIIARA----LDIQFKQGLDRMTAGSDSLKGFKDILSTKKFNKVFP 209
Query: 284 LLNQYQDIKISYTHYHFHDMNFYDYIKSFYRGGLNMYNTKYINKLIDEPCFSIDINSSYP 343
Query: 344 YVMYHEKIPTWLYFYEHYSEPTLIPTFLDDDNYFSLYKIDKDVFNDDLLIKIKSRVLRQM 403
MY +P Y P + + D + LY I + F +L K + + Sbjct: 253 SQMYSRPLP-----YGAPIVFQGKYEKDEQYPLY-IQRIRFEFEL----KEGYIPTI 299
Query: 404 XXXXXXXXXXXXXXXXXXXXXIRMIQ-DITGIDCMHIRVNSFVIYECEYFHARDIIFQNYFIK 462
                                 + ++ +T +D I+ + + +Y EY
Sbjct: 300 QIKKNPFFKGNEYLKNSGVEPVELYLTNVDLELIQEH-YELYNVEYIDGFK-----FRE 352
Query: 463 TQGKLKNKINMTSPYDYHITDDINEHPYSNEEVMLSKVVLNGLYG---
G K+ I+ + H + L+K++LN LYG +P L
Sbjct: 353 KTGLFKDFIDKWTYVKTH------EEGAKKQLAKLMLNSLYGKFASNPDVTGKVPYL 403
Query: 512 RSHFNL-FRLDDNNELYNIINGYKNTERNILFSTFVTSRSLYNLLVPFQYLTESEIDDNF 570
+ +L FR+ D YK+ + F+T+ + + Q D
Sbjct: 404 KDDGSLGFRVGDEE-----YKDPVYTPM-GVFITAWARFTTITAAQACY----DRI 449
Query: 571 IYCDTDSLYMKSVVKPLLNPSLFDPIALGKWDIENEQIDKMFVLNHKK-----YAYEVNG 625
IYCDTDS+++ P + DP LG W E+ + L K Y EV+G
Sbjct: 450 IYCDTDSIHLTGTEVPEIIKDIVDPKKLGYWAHES-TFKRAKYLRQKTYIQDIYVKEVDG 508
Query: 626 KIKIAS 631
            K+K S
Sbjct: 509 KLKECS 514
>gi|1072656|pir||S51275 DNA polymerase - phage CP-1
             >gi|836593|emb|CAA87725.1| (247794) DNA polymerase
             [Bacteriophage CP-1]
            Length = 568
 Score = 53.5 bits (126), Expect = 6e-06
 Identities = 104/464 (22%), Positives = 169/464 (36%), Gaps = 66/464 (14%)
Query: 230 LTPEQLTYIHNDVIIL--GMCHIHYSDIFPNFDYNKLTFSLNIMESYLNNEMTRFQLLNQ 287
+ PE + YIH DV IL G+ ++Y + F Y + +L + +F+
Sbjct: 152 IKPEWIDYIHVDVAILARGIFAMYYEENFTK--YTSASEALTEFKRIFRKSKRKFRDFFP 209
Query: 288 YQDIKISYTHYHFHDMNFYDYIKSFYRGGLMMYNTKYINKLIDEPCFSIDINSSYPYVMY 347

D K+ D+ G + K+ ++++ DINS YP M

Sbjct: 210 ILDEKVD------DFCRKHIVGAGRLPTLKHRGRTLNQLIDIYDINSMYPATML 257
                                                                                             Query: 348 HEKIPTWLYFYEHYSEPTLIPTFLDDDNYFSLY-KIDKDVFNDDL-LIKIKSRVLRQMXX 405 -
+P + + Y P + +D+Y+ + K D D+ L I+IK ++
Sbjct: 258 QNALPIGIP--KRYKGK---PKEIKEDHYYIYHIKADFDLKRGYLPTIQIKKKLDALRIG 312
Query: 406 XXXXXXXXXXXXXXXXXXIRMIQDITGIDCMHIRVNSFVIYECEYFHARDIIFQNYFIKTQG 465
L + + H + E F +F +Y
Sbjct: 313 VRTSDYVTTSKNEVIDLYLTNFDLDLFLKHYDATIMYVETLE-FQTESDLFDDYI---- 366
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WO 00/32825 PCT/1B99/02040

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Query: 466 KLKNKINMTSPYDYHITDDINEHPYSNEEVMLSKVVLNGLYGIPALR--SHFNLFRLDDN 523
+ Y Y E+ S E +K++LN LYG + S L LDD
Sbjct: 367 -----TTYRYK-----KENAQSPAEKQKAKIMLNSLYGKFGAKIISVKKLAYLDDK 412
Query: 524 NELYNIINGYKNTERNIL-----FSTFVTSRSLYNLLVPFQYLTESEIDDNFIYCDTDS 577
L +KN + + + FVTS + + + + Q E DNF+Y DTDS

Sbjct: 413 GILR----FKNDDEEEVQPVYAPVALFVTSIARHFIISNAQ----ENYDNFLYADTDS 462
Query: 578 LYMKSVVKPLLNPSLFDPIALGKWDIENEQIDKMFVLNHKKYAYEVNGKIKIASAGIPKN 637
L++ +L+ DP GKW E + K L K Y E+ + + K
Sbjct: 463 LHLFHSDSLVLD---IDPSEFGKWAHEGRAV-KAKYLRSKLYIEELIQEDGTTHLDV-KG 517
Query: 638 AFDTSVDFETFVREQFFDGAILENNKSIYNEQGTISIYPSKTEI 681
            ATEFFGAE++ +G IY+ +I
Sbict: 518 AGMTPEIKEKITFENFVIGATFEGKRASKQIKGGTLIYETTFKI 561
>gi|1429230|emb|CAA67649| (X99260) DNA polymerase [Bacteriophage
            B1031
            Length = 572
 Score = 49.2 bits (115), Expect = 1e-04
 Identities = 93/422 (22%), Positives = 155/422 (36%), Gaps = 88/422 (20%)
Query: 229 KLTPEQLTYIHNDVIILGMCHIHYSDIFPNFDYNKLTFSLNIMESYLNNEMTR-----FQ 283
++TPE+ YI ND+ I+ DI +++T + ++ + T+ F
Sbjct: 154 EITPEEYEYIKNDIEIIARA----LDIQFKQGLDRMTAGSDSLKGFKDILSTKKFNKVFP 209
 Query: 284 LLNQYQDIKISYTHYHFHDMNFYDYIKSFYRGGLNMYNTKYINKLIDEPCFSIDINSSYP 343
Query: 344 YVMYHEKIPTWLYFYEHYSEPTLIPTFLDDDNYFSLYKIDKDVFNDDLLIKIKSRVLRQM 403
MY +P Y P + + D + LY I + F +L K + + Sbjct: 253 SQMYSRPLP-----YGAPIVFQGKYEKDEQYPLY-IQRIRFEFEL----KEGYIPTI 299
 Query: 404 XXXXXXXXXXXXXXXXXXXXXIRMIQ-DITGIDCMHIRVNSFVIYECEYFHARDIIFQNYFIK 462
                                   ++ +T +D I+ + +Y EY
 Sbjct: 300 QIKKNPFFKGNEYLKNSGAEPVELYLTNVDLELIQEH-YEMYNVEYIDGFK-----FRE 352
 Query: 463 TQGKLKNKINMTSPYDYHITDDINEHPYSNEEVMLSKVVLNGLYG------IPAL 511
G K I+ + H + L+K++ LYG +P L
Sbjct: 353 KTGLFKEFIDKWTYVKTH-----EKGAKKQLAKLMFDSLYGKFASNPDVTGKVPYL 403
 Query: 512 RSHFNL-FRLDDNNELYNIINGYKNTERNILFSTFVTSRSLYNLLVPFQYLTESEIDDNF 570
+ +L FR+ D YK+ + F+T+ + + Q D

Sbjct: 404 KEDGSLGFRVGDEE-----YKDPVYTPM-GVFITAWARFTTITAAQACY----DRI 449
 Query: 571 IYCDTDSLYMKSVVKPLLNPSLFDPIALGKWDIENEQIDKMFVLNHKK----YAYEVNG 625
 IYCDTDS+++ P + + DP LG W E+ + L K YA EV+G
Sbjct: 450 IYCDTDSIHLTGTEVPEIIKDIVDPKKLGYWAHES-TFKRAKYLRQKTYIQDIYAKEVDG 508
 Query: 626 KI 627
 Sbjct: 509 KL 510
 >gi|1572479|emb|CAA65712| (X96987) DNA polymerase (Bacteriophage
             GA-1)
             Length = 578
  Score = 46.1 bits (107), Expect = 0.001
  Identities = 80/376 (21%), Positives = 146/376 (38%), Gaps = 54/376 (14%)
 Query: 234 QLTYIHNDVIILGMCHIHYSDIFPNFDYNKLTFSLNIMESYLNNEMTRFQLLNQYQDIKI 293
 ++ Y+ +D++I+ + +F N D+ +T + + +Y EM + +Y +

Sbjct: 162 EIEYLKHDLLIVALA---LRSMFDN-DFTSMTVGSDALNTY--KEMLGVKQWEKYFPVL- 214
 Query: 294 SYTHYHFHDMNFYDYIKSFYRGGLNMYNTKYINKLIDEPCFSIDINSSYPYVMYHEKIPT 353
                            I+ Y+GG N KY + + D+NS YP +M ++ +P
 Sbjct: 215 -----SLKVNSEIRKAYKGGFTWVNPKYQGETVYGGMV-FDVNSMYPAMMKNKLLP- 264
 Query: 354 WLYFYEHYSEPTLIPTFLDDDNYFSLYKIDKDVFNDDLLIKIKSRVLRQMXXXXXXXXX 413
                                 + + LY
                                               F + KI
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WO 00/32825 PCT/IB99/02040

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Sbjct: 265 -----YGEPVMFKGEYKKNVEYPLYIQQVRCFFELKKDKIPCIQIKGNARFGQNEYLS 317
Query: 414 XXXXXXXXLRMIQDITGIDCMHIRVNSFVIYECEYFHARDIIFQNYFIKTQGKLKNKINM 473
L +T +D I+ + + I+E E+ +F+ + I

Sbjct: 318 TSGDEYVDLY----VTNVDWELIKKH-YDIFEEEFIGG--FMFKGF------IGF 359
Query: 474 TSPYDYHITDDINEHPYSNEEVMLSKVVLNGLYGIPALRSHFN--LFRLDDNNELYNIIN 531
Y + N S E+ + +K++LN LYG A + LD+N L
Sbjct: 360 FDEYIDRFMEIKNSPDSSAEQSLQAKIMLNSLYGKFATNPDITGKVPYLDENGVLKFRKG 419
Query: 532 GYKNTERNILFST---FVTSRSLYNLLVPFQYLTESEIDDNFIYCDTDSLYMKSVVKPLL 588
K ER+ +++ F+T+ + N+L Q L FIY DTDS++++ + +
Sbjct: 420 ELK--ERDPVYTPMGCFITAYARENILSNAQKLYP-----RFIYADTDSIHVEGLGEVDA 472
Query: 589 NPSLFDPIALGKWDIE 604
               + DP LG WD E
Sbjct: 473 IKDVIDPKKLGYWDHE 488
>gi|118851|sp|P06950|DPOL_BPPZA DNA POLYMERASE (EARLY PROTEIN GP2)
>gi|75812|pir||ERBP2Z DNA-directed DNA polymerase (EC
            2.7.7.7) - phage PZA >gi|216051 (M11813) gene 2 product [Bacteriophage PZA] >gi|224741|prf||1112171E ORF 2
            [Bacteriophage PZA]
            Length = 572
 Score = 45.3 bits (105), Expect = 0.002
 Identities = 98/461 (21%), Positives = 166/461 (35%), Gaps = 110/461 (23%)
Query: 198 QLKTDFNYTIFDKDNDMNDSEAYDYAVKCFAKLTPEQLTYIHNDVIILGMCHIHYSDIFP 257
++ DF T+ D D + Y ++TP++ YI ND+ I+ + I
Sbjct: 129 KIAKDFKLTVLKGDIDYHKERPVGY-----EITPDEYAYIKNDIQIIAEALL----IQF 178
Query: 258 NFDYNKLTFSLNIMESYLNNEMTR-----FQLLNQYQDIKISYTHYHFHDMNFYDYIKSF 312
Sbjct: 179 KQGLDRMTAGSDDLKGFKDIITTKKFKKVFPTLSLGLDKEVRYA----- 222
Query: 313 YRGGLNMYNTKYINKLIDEPCFSIDINSSYPYVMYHEKIPTWLYFYEHYSEPTLIPT--F 370
            YRGG N ++ K I E D+NS YP MY +P
Sbjct: 223 YRGGFTWLNDRFKEKEIGEGMV-FDVNSLYPAQMYSRLLP------YGEPIVFEGKYV 273
Sbjct: 274 WDEDYPLHIQHIRCEFELKEGYIPTIQIK-RSRFYKGNEYLKSSGGEIADLW------ 324
 Query: 426 QDITGIDCMHIRVNSFVIYECEYFHARDIIFQNYFIKTQGKLKNKINMTSPYDYHITDDI 485
++ +D + + + +Y EY F T G K+ I+ + I
Sbjct: 325 --VSNVD-LELMKEHYDLYNVEYISGLK-----FKATTGLFKDFIDKWTHIKTTSEGAI 375
Query: 486 NEHPYSNEEVMLSKVVLNGLYG-------IPALRSHFNL-FRLDDNNELYNIINGY 533
+ L+K++LN LYG +P L+ + L FRL G
Sbjct: 376 KQ------LAKLMLNSLYGKFASNPDVTGKVPYLKENGALGFRL------GE 415
 Query: 534 KNTERNIL--FSTFVTSRSLYNLLVPFQYLTESEIDDNFIYCDTDSLYMKSVVKPLLNPS 591
                         F+T+ + Y + Q D IYCDTDS+++
 Sbjct: 416 EETKDPVYTPMGVFITAWARYTTITAAQACF----DRIIYCDTDSIHLTGTEIPDVIKD 470
 Query: 592 LFDPIALGKWDIENEQIDKMFVLNHKKYAY----EVNGKI 627
             + DP LG W E+ + L KY
 Sbjct: 471 IVDPKKLGYWAHES-TFKRAKYLRQKTYIQDIYMKEVDGKL 510
 >gi|2435429 (AF012250) unassigned reading frame (possible DNA
            polymerase) [Physarum polycephalum]
             Length = 544
  Score = 44.9 bits (104), Expect = 0.002
  Identities = 118/545 (21%), Positives = 206/545 (37%), Gaps = 104/545 (19%)
 Query: 179 TSIATLGKKLLDGGYLTESQLKTDFNYTIFDKDNDMNDSEAYDYAVKCFAKLTPEQLTYI 238 T + L K L D + T Q F N M Y + CF L P++ I
 T + L K L D + T Q F N M Y + CF L P++ I
Sbjct: 62 TQLFNLLKSLQDSSFYTFKQ------FTYQNIM----YSLEISCF--LYPKKKILI 105
 Query: 239 HNDVIILGMCHIHYSDIFPNFD----YNKL--TFSLNIMESY-LNNEMTRFQLLNQYQD 290
                     +I Y+D+ ++ YN++ +++NI Y L+
 Sbjct: 106 -KDLYNFFSENIIYNDVVKDYKLLAILYNEIQTAYNININRKYILSTASLSLRIFKKSFP 164
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WO 00/32825 PCT/IB99/02040

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Query: 291 IKISYTHYHFHDMNFYDYIKSFYRGGLNMYNTKYINKLIDEPCFSIDINSSYPYVMYHEK 350

K + D + +YI+ Y GG N I + + + + D+NS YPY+M EK

Sbjct: 165 EKYRLIPHLTRDED--NYIRKSYIGGRNE-----IFEHVAQRNYFYDVNSLYPYIMKKEK 217
Query: 351 IPTWLYFYEHYSEPTLIPTFLDD-DNYFS----LYKIDKDVFNDDLL---IKIKSRVLRQ 402
+P + Y + + F + +N+F L I+K N +L + IK+ V
Sbjct: 218 MPIGI---PEYRDKEYMKKFEKNIENFFGFIDVLITIEKTNNNIPVLPYRMGIKNNV-EV 273
Query: 403 MXXXXXXXXXXXXXXXXXXXXXXXIRMIQDITGIDCMHIRVNSFVIYECEYFHARDIIFQNYFIK 462
L + Q I+ IY + ++++F+ Y +
Sbjct: 274 GIIYAKGTLRGIYFSEEIKLALKQGYKIIE------IYSAYEYKEKEVVFEEYVEQ 323
Query: 463 TQGK-LKNKINMTSPYDYHITDDINEHPYSNEEVMLSKVVLNGLYG-----IPALRS 513
+ LK K D + D L K + LN LYG I +
Sbjct: 324 MYNRRLKAK-----DPALKD-------LYKKLLNTLYGRFGLVYEQIDIISP 363
Query: 514 HFNLFRLDDNNELYNIINGYKNTERNILFSTFVTSRSLYNLLVPFQYLTESEIDDNFIYC 573
L + DN + + + + + N ++ + + + + F Y T + + IY
Sbjct: 364 EKEL--ITDNTYISHDTTEFIDITANTCYNNIAITSAITSYARIFMYNTILNYNLHVIYI 421
Query: 574 DTDSLYMKSVVKPLLNPSLFDPIALGKWDIENEQIDKMFVLMHKKYAY-EVNGKIKIASA 632
DTD L++K+ P+ +L +GK+ +E+ + F+ N K Y Y +N I
Sbjct: 422 DTDGLFLKN---PIPDIALTTSKEMGKFRLESINAEAHFIAN-KFYIYAPINSPIIYKFK 477
Query: 633 GIPK-----NAFDTSVDFETFVR----EQFFDGAIIENNKSIYNEQGT-----ISIYPSK 678
                     N D + + +F +I NN Y+Q+ I Y+
Sbict: 478 GIPLOKPIFNIHDIITOHKKILNITLGHHYFTFSIRLMNNQTYSFQASRKRKLIPNYKTT 537
Query: 679 TEIVC 683
                I+C
Sbict: 538 PWIIC 542
>qi|1084487|pir||S41618 DNA polymerase - slime mold (Physarum
              polycephalum) >gi|509721|dbj|BAA06121.1| (D29637) DNA
              polymerase [Physarum polycephalum]
              Length = 547
 Score = 44.9 bits (104), Expect = 0.002
 Identities = 118/545 (21%), Positives = 206/545 (37%), Gaps = 104/545 (19%)
Query: 179 TSIATLGKKLLDGGYLTESQLKTDFNYTIFDKDNDMNDSEAYDYAVKCFAKLTPEQLTYI 238
T + L K L D + T Q F N M Y + CF L P++ I

Sbjct: 65 TQLFNLLKSLQDSSFYTFKQ------FTYQNIM-----YSLEISCF--LYPKKKILI 108
Query: 239 HNDVIILGMCHIHYSDIFPNFD-----YNKL--TFSLNIMESY-LNNEMTRFQLLNQYQD 290
                        +I Y+D+ ++ YN++ +++NI Y L+
Sbjct: 109 -KDLYNFFSENIIYNDVVKDYKLLAILYNEIQTAYNININRKYILSTASLSLRIFKKSFP 167
Query: 291 IKISYTHYHFHDMNFYDYIKSFYRGGLNMYNTKYINKLIDEPCFSIDINSSYPYVMYHEK 350
K + D + +YI+ Y GG N I + + + D+NS YPY+M EK
Sbjct: 168 EKYRLIPHLTRDED-NYIRKSYIGGRNE----IFEHVAQRNYFYDVNSLYPYIMKKEK 220
Query: 351 IPTWLYFYEHYSEPTLIPTFLDD-DNYFS----LYKIDKDVFNDDLL---IKIKSRVLRQ 402 +P + Y + + F + +N+F L I+K N +L + IK+ V
Sbjct: 221 MPIGI---PEYRDKEYMKKFEKNIENFFGFIDVLITIEKTNNNIPVLPYRMGIKNNV-EV 276
Query: 403 MXXXXXXXXXXXXXXXXXXXXIRMIQDITGIDCMHIRVNSFVIYECEYFHARDIIFQNYFIK 462
L + Q I+ IY + ++++F+ Y +
Sbjct: 277 GIIYAKGTLRGIYFSEEIKLALKQGYKIIE------IYSAYEYKEKEVVFEEYVEQ 326
Query: 463 TQGK-LKNKINMTSPYDYHITDDINEHPYSNEEVMLSKVVLNGLYG------IPALRS 513
+ LK K D + D L K + LN LYG I +
Sbjct: 327 MYNRRLKAK-----DPALKD--------LYKKLLNTLYGRFGLVYEQIDIISP 366
Query: 514 HFNLFRLDDNNELYNIINGYKNTERNILFSTFVTSRSLYNLLVPFQYLTESEIDDNFIYC 573

L + DN + + + + + + + + + F Y T + + IY

Sbjct: 367 EKEL--ITDNTYISHDTTEFIDITANTCYNNIAITSAITSYARIFMYNTILNYNLHVIYI 424
Query: 574 DTDSLYMKSVVKPLLNPSLFDPIALGKWDIENEQIDKMFVLNHKKYAY-EVNGKIKIASA 632
DTD L++K+ P+ +L +GK+ +E+ + F+ N K Y Y +N I
Sbjct: 425 DTDGLFLKN---PIPDIALTTSKEMGKFRLESINAEAHFIAN-KFYIYAPINSPIIYKFK 480
Ouery: 633 GIPK-----NAFDTSVDFETFVR----EQFFDGAILENNKSIYNEQGT-----ISIYPSK 678
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+F +I NN Y+Q+ I Y+
            GIP
                      N D
Sbjct: 481 GIPLQKPIFNIHDIITQHKKILNITLGHHYFTFSIRLNNNQTYSFQASRKRKLIPNYKTT 540
Query: 679 TEIVC 683
Sbjct: 541 PWIIC 545
>gi|4877819|gb|AAD31446.1| (AF133505) DNA polymerase [Neurospora
            crassa]
            Length = 1035
 Score = 44.1 bits (102), Expect = 0.004
 Identities = 36/172 (20%), Positives = 82/172 (46%), Gaps = 14/172 (8%)
Query: 521 DDNNELYNIINGYKNTERNILFSTFVTSRSLYNLLVPFQYLTESEIDDNFIYCDTDSLYM 580
+ N EL + ++G K+ I ++ ++ +++ +++ S Y DTDS+++
Sbjct: 817 EKNYELLSYLDGEKDDGFIINSTSIAAATASWSRILMYKHIINSA------YTDTDSIFV 870
Query: 581 KSVVKPLLNPSLFDPIALGKWDIENEQIDKMFVLNHKKYAYEVNGKIKIASAGIPKNAFD 640
                              K + + I + ++ K Y + GK++I GI KN +
Sbjct: 871 E---KPLDSAFIGEGCGKFKAEYNGQLIKRAIFISGKLYLLDFGGKLEIKCKGITKNKDN 927
Query: 641 TSVDFETFVREQFFDG---AIIENNKSIYNEQGTISIYPSKTEIVCGNVYDE 689
                     E ++G + +
                                            E GT+++ K ++ G YD+
Sbjct: 928 TTHNLDINDFEALYNGESRVLFQERWGRSLELGTVTVKYQKYNLISG--YDK 977
>gi|461962|sp|P33537|DPOM_NEUCR PROBABLE DNA POLYMERASE
            >gi|283351|pir||S26985 probable DNA-directed DNA
polymerase (EC 2.7.7.7) - Neurospora crassa
            mitochondrion plasmid maranhar (SGC3)
            >gi|578156|emb|CAA39046| (X55361) putative DNA
            polymerase [Neurospora crassa]
            Length = 1021
 Score = 44.1 bits (102), Expect = 0.004
 Identities = 36/172 (20%), Positives = 82/172 (46%), Gaps = 14/172 (8%)
Query: 521 DDNNELYNIINGYKNTERNILFSTFVTSRSLYNLLVPFQYLTESEIDDNFIYCDTDSLYM 580
+ N EL + ++G K+ I ++ ++ ++ +++ S Y DTDS+++
Sbjct: 815 EKNYELLSYLDGEKDDGFIINSTSIAAATASWSRILMYKHIINSA------YTDTDSIFV 868
                                                                   Y DTDS+++
Query: 581 KSVVKPLLNPSLFDPIALGKWDIENEQIDKMFVLNHKKYAYEVNGKIKIASAGIPKNAFD 640
+ KPL + + + K + + I + + + K Y + GK++I GI KN + Sbjct: 869 E---KPLDSAFIGEGCGKFKAEYNGQLIKRAIFISGKLYLLDFGGKLEIKCKGITKNKDN 925
Query: 641 TSVDFETFVREQFFDG---AIIENNKSIYNEQGTISIYPSKTEIVCGNVYDE 689
T+ + + E ++G + + E GT+++ K ++ G YD+
Sbjct: 926 TTHNLDINDFEALYNGESRVLFQERWGRSLELGTVTVKYQKYNLISG--YDK 975
>gi|2499511|sp|Q12471|6P22_YEAST 6-PHOSPHOFRUCTO-2-KINASE 2
             (PHOSPHOFRUCTOKINASE 2 II) (6PF-2-K 2)
             >gi|2131162|pir||S61066 6-phosphofructo-2-kinase (EC
            2.7.1.105) - yeast (Saccharomyces cerevisiae)
>gi|2131163|pir||571026 6-phosphofructo-2-kinase (EC
             2.7.1.105) - yeast (Saccharomyces cerevisiae)
             >gi|1085116|emb|CAA62371| (X90861)
             6-phosphofructo-2-kinase [Saccharomyces cerevisiae]
             >gi|1420028|emb|CAA99157| (274878) ORF YOL136c
[Saccharomyces cerevisiae] >gi|1628439|emb|CAA64733|
             (X95465) 6-phosphofructo-2-kinase [Saccharomyces
             cerevisiae)
             Length = 397
  Score = 40.6 bits (93), Expect = 0.041
  Identities = 48/208 (23%), Positives = 92/208 (44%), Gaps = 29/208 (13%)
                                                                                          ____
 Query: 175 MKTNTSIATLGKKLLDGGYLTESQLKTDFNYTIFDKDNDMNDSEAYDYAVKCFAKLTPEQ 234
++ S AT+ K LL L+ + + FN K+ND ++ +A++T ++
Sbjct: 139 IRRQISCATISKPLL---LSNTSSEDLFN---PKNNDKKET-------YARITLQK 181
 Query: 235 LTY-IHNDVIILGMCHIHYSDIFPNFDYNKLTFSLNIMESYLNNEMTRFQLLN----QYQD 290
L + I+ND +G+ S I + F + S+ +E++ F L+ Q

Sbjct: 182 LFHEINNDECDVGIFDATNSTI-----ERRRFIFEEVCSFNTDELSSFNLVPIILQVSC 235
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Query: 291 IKISYTHYHFHDMNFY-DYIKSFYRGGLNMYNTKYINKLIDEPCFSID-INSSYPYVMYH 348
S+ Y+ H+ +F DY+ Y + + + + FS+D N + Y+ H
Sbjct: 236 FNRSFIKYNIHNKSFNEDYLDKPYELAIKDFAKRLKHYYSQFTPFSLDEFNQIHRYISQH 295
Query: 349 EKIPTWLYFYEHYSEPTLIPTFLDDDNY 376
E+I T L+F+ + + P L+ +Y
Sbjct: 296 EEIDTSLFFFNVINAGVVEPHSLNQSHY 323
>gi|2258375|gb|AAD11909.1| (AP007261) transcription initiation
            factor sigma (Reclinomonas americana)
            Length = 532
 Score = 39.9 bits (91), Expect = 0.070
 Identities = 49/205 (23%), Positives = 84/205 (40%), Gaps = 14/205 (6%)
Query: 100 NHFLLKDTMRYFDNITRENIYLKSAEENEHTLKMKEATILAKNQNVIL---EKRVKSSIN 156
N+ + + F + ++IY+ + +KE L K NVI+ K +K N
Sbjct: 177 NYLVKNSYLNLFKTVPHDSIYMNYSYIQTPLNILKEYLQLIKIINVIILQINKNIKKKNN 236
Query: 157 LDLTMFLNGFKFNIIDNFM---KTNTSIATLGKKLLDGGYLTESQLKTDFNYTIFDKDND 213
            L++++FL F + N++ K + + + K L Y+T L T Y
Sbjct: 237 LNISLFLYKFYQELKWNYIFINKISRNTQKINIKTLKNSYITFYNLITFIQYYTTKKQRL 296
Query: 214 MNDSEAYDYAVKCFAK--LTPEQLTYIHNDVIILGMCHIHYSDIFPNFDYN-KLTFSLNI 270
                      +K F K P+ +N +I G+ HI+ + N K+T I
Sbjct: 297 KKDIFYKQIFIKTFLKQHKIPKINKIKNNSLIKYGLTHIYDMILISILRENIKVTLKNRI 356
Query: 271 MESYLNNEMTRFQLLNQYQDIKISY 295
+ +Y+ T + QY +KI Y
Sbjct: 357 IFNYMPYITT---ISKQY--VKIGY 376
>gi|15734|emb|CAA37450| (X53370) DNA polymerase (AA 1-575)
            [Bacteriophage phi-29]
            Length = 575
 Score = 39.5 bits (90), Expect = 0.092 Identities = 41/150 (27%), Positives = 64/150 (42%), Gaps = 36/150 (24%)
Query: 497 LSKVVLNGLYG------IPALRSHFNL-FRLDDNNELYNIINGYKNTERNIL--F 542
            L+K++LN LYG +P L+ + L FRL
Sbjct: 381 LAKLMLNSLYGKFASNPDVTGKVPYLKENGALGFRL------GEEETKDPVYTPM 429
Query: 543 STFVTSRSLYNLLVPFQYLTESEIDDNFIYCDTDSLYMKSVVKPLLNPSLFDPIALGKWD 602
F+T+ + Y + Q D IYCDTDS+++ P + + DP LG W
Sbjct: 430 GVFITAWARYTTITAAQACY----DRIIYCDTDSIHLTGTEIPDVIKDIVDPKKLGYWA 484
Query: 603 IENEQIDKMFVLNHKKYAY----EVNGKI 627
Sbjct: 485 HES-TFKRVKYLRQKTYIQDIYMKEVDGKL 513
Query= pt | 110872 44AHJDORF002 Phage 44AHJD ORF | 3789-5732 | 3 1
          (647 letters)
>gi|135273|sp|P27622|TAGC_BACSU TEICHOIC ACID BIOSYNTHESIS PROTEIN C
            >gi|478126|pir||D49757 techoic acid biosynthesis protein
            tagC - Bacillus subtilis (strain 168) >gi|143727
            (M57497) putative (Bacillus subtilis)
            >gi|2636103|emb|CAB15594.1| (299122) alternate gene
            name: dinC (Bacillus subtilis)
            Length = 442
 Score = 112 bits (278), Expect = 7e-24 Identities = 91/314 (28%), Positives = 147/314 (45%), Gaps = 58/314 (18%)
Query: 152 FELNELEPKFVMGFGGIRNAVNQSINIDKETNHMYSTQSDS----QKPEGFWINKLTPSG 207
                             V QS N D++ + +Y+TQ S
Sbjct: 7 FDFTNITPKLFTELRVADKTVLQSFNFDEKNHQIYTTQVASGLGKDNTQSYRITRLSLEG 66
Query: 208 DLISSMRIVQGGHGTTIGLERQSNGEMKIWLHHD-----GVAKLLQVAYKDNYVLDLEEA 262
              + SM + GGHGT IG+E + NG + IW +D
Sbjct: 67 LQLDSMLLKHGGHGTNIGIENR-NGTIYIWSLYDKPNETDKSELVCFPYKAGATLD-ENS 124
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Query: 263 KGLTDYTPQSLLNKHTFTPLIDEANDKLILRFGDGTIQVRSRADVKNHIDNVEKEMTIDN 322
          K L ++ H TP +D N +L +R + D KN+ N ++ +TI N
Sbjct: 125 KELQRFSNMPF--DHRVTPALDMKNRQLAIR------QYDTKNN--NNKQWVTIFN 170
Query: 323 SE----NNDN-------RWMQGIAVDGDDLYWLSGNSSVNSHVQIGKYSLTTGQKI 367
                               ++QG +D LYW +G+++ S+ +
              N +N
Sbjct: 171 LDDAIANKNNPLYTINIPDELHYLQGFFLDDGYLYWYTGDTNSKSYPNL-----ITV 222
Query: 368 YDYPFKLSYQDGINFPRD-----NFKEPEGICIYTNPKTKRKSLLLAMTNGGGGKRFH 420
          +D K+ Q I +D NF+EPEGIC+YTNP+T KSL++ +T+G G R
Sbjct: 223 FDSDNKIVLOKEITVGKDLSTRYENNFREPEGICMYTNPETGAKSLMVGITSGKEGNRIS 282
Ouery: 421 NLYGFFQLGEYEHF 434
                  YE+F
Sbjct: 283 RIYAYH---SYENF 293
>gi|142847 (M64050) DNase inhibitor (Bacillus subtilis)
Score = 51.9 bits (122), Expect = 1e-05
Identities = 35/116 (30%), Positives = 55/116 (47%), Gaps = 10/116 (8%)
Query: 152 FELNELEPKFVMGFGGIRNAVNQSINIDKETNHMYSTQSDS----QKPEGFWINKLTPSG 207
                             V QS N D++ + +Y+TQ S + + I +L+ G
          F+ + PK
Sbjct: 7 FDFTNITPKLFTELRVADKTVLQSFNFDEKNHQIYTTQVASGLGKDNTQSYRITRLSLEG 66
Query: 208 DLISSMRIVQGGHGTTIGLERQSNGEMKIWLHHD-----GVAKLLQVAYKDNYVLD 258
            + SM + GGHGT IG+E + NG + IW +D ++L+ YK LD
Sbjct: 67 LQLDSMLLKHGGHGTNIGMENR-NGTIYIWSLYDKPNETDKSELVCFPYKAGATLD 121
>gi|4038407 (AF103943) factor C protein precursor (Streptomyces
          griseus]
          Length = 324
 Score = 39.1 bits (89), Expect = 0.10
 Identities = 61/269 (22%), Positives = 102/269 (37%), Gaps = 33/269 (12%)
Query: 172 VNQSINIDKETNHMYSTQSDSQKPEG---FWINKLTPSGDLISSMRIVQGGHGTTIGLER 228
V QS D ++ Q S P+ I +L SG+ + M ++ GHG +IG +
Sbjct: 66 VQQSFTFDIVNRRLFVAQLKSGSPDDSGDLCITQLDFSGNKLGHMYLLGFGHGVSIGAQ- 124
Query: 229 QSNGEMKIWLHHDGVAKLLQVAYKDNYVLDLEEAKGLTDYTPQSLLNKHTFTP------ 281
                                 + + + G T
                                                     SLKH
               + +W D +
Sbjct: 125 PVGADTYLWTEVD-----VNSNARGTRLARFKWNNGATLSRTSSALAKHQPVPGATEMTC 179
Query: 282 LIDEANDKLILRFGDGTIQVRSRADVKNHIDNVEKEMTIDNSENNDNRWMQGIAVDGDDL 341
ID N+++ +R+ + + + +V + V + D QG A+ G + Sbjct: 180 AIDPVNNRMAIRYLTASGRRYGIYNVADIAAGVYDKPLSDVPHPTGLGTFQGYALYGSYV 239
Query: 342 YWLSGN-----SSVNSHVQIGKYSLTTGQKIYDYPFKLSYQDGINFPRDNFKEPEGIC 394
          Y L+GN + NS+V + TG + + + G
Sbjct: 240 YQLTGNPYGPDNPNPGNSYVS--SVDVNTGALVQ----RAFTRAGSTL---TFREPEGMG 290
Query: 395 IYTNPKTKRKSLLLAMTNGGGGKRFHNLY 423
                + + L L +G G R NL+
          IY
Sbjct: 291 IYRTAAGEVR-LFLGFASGVAGDRRSNLF 318
Query= pt|110873 44AHJDORF003 Phage 44AHJD ORF |6626-8389|2 1
         (587 letters)
>gi|138123|sp|P04331|VG9_BPPH2 TAIL PROTEIN (LATE PROTEIN GP9)
          >gi|75850|pir||WMBPT9 gene 9 protein - phage phi-29
           >gi|215327 (M14782) tail protein [Bacteriophage phi-29]
           >gi|225364|prf||1301270D gene 9 [Bacillus sp.]
           Length = 599
                                                                            - 10 40 50
- 20 40
- 20 40
 Score = 92.4 bits (226), Expect = 8e-18
 Identities = 126/618 (20%), Positives = 251/618 (40%), Gaps = 71/618 (11%)
          {\tt TNFKFFYNTPFT-DYQNTIHFNSNKERDDYFLNGRHFKSLDYSKQPY-NFIRDRMEINVD~62}
         TN + + PF+ DY+NT F S+ + ++F R + + SK + F ++ ++V
TNVRILADVPFSNDYKNTRWFTSSSNQYNWF--NRKSRVYEMSKVTFMGFRENKPYVSVS 66
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Query: 63 MQWHDAQGINYMTFLS-DFEDRRYYAFVNQIEYVNDVVVKIYFVIDTIMTYTQGNVLEQL 121
                   +Y+ F + D+ ++ +YAFV ++E+ N V ++F ID + T+ ++
Sbjct: 67 LPIDKLYSASYIMFQNADYGNKWFYAFVTELEFKNSAVTYVHFEIDVLQTWMFDMKFQES 126
Query: 122 SNVNIERQHLSKRTYNYMLPMLRNNDDVLKVSNKNYVYNQMQQYLENLVLFQSSADLSKK 181
                  I R+H+ K + P + D+ L ++ ++
Sbjct: 127 F---IVREHV-KLWNDDGTPTINTIDEGLSYGSEYDIVSVENHKPYDDMMFLVIISKSIM 182
Query: 182 FGT--KKEPNLDTSKGTIYDNITSPVNLYVMEYGDFINFMDKMSAYPWITQNFQK----V 235
GT ++E L+ ++ + + P+ Y+ + + D +I N V
Sbjct: 183 HGTPGEEESRLNDINASL-NGMPQPLCYYIHPF----YKDGKVPKTYIGDNNANLSPIV 236
Query: 236 QMLPKDFINTKDLEDVKTSEKITGLKTLKQGGKSKEWSLK-DLSL-----SFSNLQ 285
               ML' F + D+ + +T LK K+ + LK D +
Sbjct: 237 NMLTNIFSQKSAVNDI-VNMYVTDYIGLKLDYKNGDKELKLDKDMFEQAGIADDKHGNVD 295
Query: 286 EMMLSK------KDEFKHMIRNEYMTIEFYDWNGNTMLLDAGKISQK 326
+ + K KD+ ++ Y E D+ GN M L I+ Sbjet: 296 TIFVKKIPDYEALEIDTGDKWGGFTKDQESKLMMYPYCVTEITDFKGNHMNLKTEYINNS 355
Query: 327 TGVKLRTKSIIGYHNEVRVYPVDYNSAENDRPILAKNKEILIDTGSFLNTNITFNSFAQV 386
+K++ + +G N+V DYN+ D + N+ S +N N
Sbjct: 356 K-LKIQVRGSLGVSNKVAYSVQDYNA---DSALSGGNRLTASLDSSLINNNPN----- 404
Query: 387 PILINNGILGQSQQANRQ--KNAESQLITNRIDNVLNG---SDPKSRFYDAVSVASNLSP 441
I I N L Q N+ +N +S ++ N I ++ G + + A+ +AS++
Sbjct: 405 DIAILNDYLSAYLQGNKNSLENQKSSILFNGIMGMIGGGISAGASAAGGSALGMASSV-- 462
Query: 442 TALFGKFNEEYNFYKQQQAEYKDLALQPPSVTESEMGNAFQLANSINGLTMKISVPSPKE 501 T + + QA+ D+A PP +T+ AF N G+ + +
Sbjct: 463 TGMTSTAGNAVLQMQAMQAKQADIANIPPQLTKMGGNTAFDYGNGYRGVYVIKKQLKAEY 522
Query: 502 ITFLQKYYMLFGFEVNDYNSFIEPINSMTVCNYLKCTGTYTIRDIDPMLMEQLKAILESG 561
L ++ +G+++N + + NY++ + DI+ +++++ I ++G
Sbjct: 523 RRSLSSFFHKYGYKINRVKK--PNLRTRKAFNYVQTKDCFISGDINNNDLQEIRTIFDNG 580
Query: 562 VRFWHNDGSGNPMLQNPL 579
              + WH D GN ++N L
Sbjct: 581 ITLWHTDNIGNYSVENEL 598
>gi|138124|sp|P07534|VG9_BPPZA TAIL PROTEIN (LATE PROTEIN GP9)
              >gi|75849|pir||WMBP9Z gene 9 protein - phage PZA
              >gi|216058 (M11813) tail protein (Bacteriophage PZA)
              Length = 599
 Score = 81.9 bits (199), Expect = 1e-14
Identities = 127/618 (20%), Positives = 248/618 (39%), Gaps = 71/618 (11%)
Query: 5 TNFKFFYNTPFT-DYQNTIHFNSNKERDDYFLNGRHFKSLDYSKQPYNFIRDRME-INVD 62
TN + PF+ DY+NT F S+ + ++F + + SK + R+ I+V
Sbjct: 9 TNVRILADVPFSNDYKNTRWFTSSSNQYNWF--NSKTRVYEMSKVTFQGFRENKSYISVS 66
Query: 63 MQWHDAQGINYMTFLS-DFEDRRYYAFVNQIEYVNDVVVKIYFVIDTIMTYTQGNVLEQL 121 ++ +Y+ F + D+ ++ +YAFV ++EY N ++F ID + T+ N+ Q
Sbjct: 67 LRLDLLYNASYIMFQNADYGNKWFYAFVTELEYKNVGTTYVHFEIDVLQTW-MFNIKFQE 125
Query: 122 SNVNIERQHLSKRTYNYMLPMLRNNDDVLKVSNKNYVYN--QMQQYLENLVLFQSSADLS 179
S I R+H+ K + P + D+ L ++ + + + Y + + L S +
Sbjct: 126 SF--IVREHV-KLWNDDGTPTINTIDEGLNYGSEYDIVSVENHRPYDDMMFLVVISKSIM 182
Query: 180 KKFGTKKEPNLDTSKGTIYDNITSPVNLYVMEY------GD------FINFMDK 221
+ E L+ ++ ++ P+ Y+ + GD +N +
Sbjet: 183 HGTAGEAESRLNDINASL-NGMPQPLCYYIHPFYKDGKVPKTFIGDNNANLSPIVNMLTN 241
Query: 222 MSAYPWITQNFQKVQMLPKDFINTK-------DLEDVKTSEKITGLKTLKQGGKSKEWS 273 + + N V M D+I K +L+ K + G+ K G + Sbjct: 242 IFSQKSAVNNI--VNMYVTDYIGLKLDYKNGDKELKLDKDMFEQAGIADDKHGNVDTIFV 299
Query: 274 LKDL---SLSFSNLQEMMLSKKDEFKHMIRNEYMTIEFYDWNGNTMLLDAGKISQKTGVK 330 K +L + KD+ ++ Y E D+ GN M L I +K Sbjet: 300 KKIPDYETLEIDTGDKWGGFTKDQESKLMMYPYCVTEVTDFKGNHMNLKTEYIDNNK-LK 358
Query: 331 LRTKSIIGYHNEVRVYPVDYNSAENDRPILAKNKEILIDTGSFLNTNITFNSFAQVPILI 390
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++ + +G N+V DYN+ + L+ +

L+T++ N+ + I+

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Sbjct: 359 IQVRGSLGVSNKVAYSIQDYNAGGS----LSGGDRLTAS----LDTSLINNNPNDIAII- 409
Query: 391 NNGILGQSQQANRQ--KNAESQLITNRIDNVLNGSDPKSRFYDAVSVASNLSP----- 441 N L Q N+ +N +S ++ N I +L G A + A SP
Sbjct: 410 -NDYLSAYLQGNKNSLENQKSSILFNGIVGMLGGG------VSAGASAVGRSPFGLASSV 462
Query: 442 TALFGKFNEEYNFYKQQQAEYKDLALQPPSVTESEMGNAFQIANSINGLTMKISVPSPKE 501 T + QA+ D+A PP +T+ AF N G+ + +
Sbjct: 463 TGMTSTAGNAVLDMQALQAKQADIANIPPQLTKMGGNTAFDYGNGYRGVYVIKKQLKAEY 522
Query: 502 ITFLQKYYMLFGFEVNDYNSFIEPINSMTVCNYLKCTGTYTIRDIDPMLMEQLKAILESG 561
                                    + + NY++ + DI+ +++++ I ++G
Sbjct: 523 RRSLSSFFHKYGYKINRVKK--PNLRTRKAYNYIQTKDCFISGDINNNDLQEIRTIFDNG 580
Query: 562 VRFWHNDGSGNPMLQNPL 579
             + WH D GN ++N L
Sbjct: 581 ITLWHTDDIGNYSVENEL 598
>gi|1429238|emb|CAA67657| (X99260) tail protein [Bacteriophage B103]
            Length = 598
 Score = 77.6 bits (188), Expect = 2e-13
Identities = 130/623 (20%), Positives = 240/623 (37%), Gaps = 86/623 (13%)
Query: 5 TNFKFFYNTPFT-DYQNTIHFNSNKERDDYFLNGRHFKSLDYSKQPYNFI---RDRMEIN 60
T+ + F N PF+ DY++T F + + YF + K + NF+ I
Sbjct: 9 TDVRIFSNVPFSNDYKSTRWFTNADAQYSYF---NAKPRVHVINBCNFVGLKEGTPHIR 64
Query: 61 VDMQWHDAQGINYMTFLS-DFEDRRYYAFVNQIEYVNDVVVKIYFVIDTIMTYTQGNVLE 119
                       YM F + + ++ +Y FV ++EYVN V +YF ID I T+
Sbjct: 65 VNKRIDDLYNACYMIFRNTQYSNKWFYCFVTRLEYVNSGVTNLYFEIDVIQTW-MFDFKF 123
Query: 120 QLSNVNIERQHLSKRTYNYMLPMLRNNDDVLKVSNKNYVYNQMQQYLENLVLFQSSADLS 179
QS+EQ+P+D+L+VQ+FFSSbjct: 124 QPSYIVREHQEMWDANNE---PLTNTIDEGLNYGTEYDVVAVEQYKPYGDLMFMVCISKS 180
Query: 180 KKFGTKKEPNLDTSKGTIYDNITS---PVNLYVMEYGDFINFMDKMSAYPWITQNFQKVQ 236
            K T E G I NI P++ YV + + D S P +T +VQ
Sbjct: 181 KMHATAGET---PKAGEIAANINGAPQPLSYYVHPF-----YEDGSS--PKVTIGSNEVQ 230
Query: 237 ML-PKDFINTKDLEDVKTSEKITGLKT-----LKQGGKSKEWSLKDLSLSFSNL----- 284
+ P DF+ ++ + ++ T + +K SL+D ++ Sbjct: 231 VSKPTDFLKNMFTQEHAVNNIVSLYVTDYIGLNIHYDESAKTMSLRDTMFEHAQIADDKH 290
Query: 285 -----DWNGNTMLLDAGK 322
+E + +F NE + Y D+ GN + +
Sbjct: 291 PNVNTIYLKEVKEYEEKTIDTGYKFASFANNEQSKLLMYPYCVTTITDFKGNQIDIKNEY 350
Query: 323 ISQKTGVKLRTKSIIGYHNEVRVYPVDYNS---AENDRPILAKNKEILIDTGSFLNTNIT 379
++ + + K++ + +G N+V DYN+ D+ +A NT++
Sbjct: 351 VNG-SNLKIQVRGSLGVSNKVTYSVQDYNADTTLSGDQNLTAS-----CNTSLI 398
Query: 380 FNSFAQVPILINNGILGQSQQANRQ--KNAESQLITNRIDNVLN---GSDPKSRFYDAVS 434 N+ V I+ N L Q N+ +N + ++ N + ++L G+ + AV
Sbjct: 399 NNNPNDVAII--NDYLSAYLQGNKNSLENQKDSILFNGVMSMLGNGIGAVGSAATGSAVG 456
Query: 435 VASNLSPTALFGKFNEEYNFYKQQQAEYKDLALQPPSVTESEMGNAFQIANSINGLTMKI 494 VAS S T + + QA+ D+A PP + + A+ N G+ +
Sbjct: 457 VAS--SATGMVSSAGNAVLQIQGMQAKQADIANTPPQLVKMGGNTAYDYGNGYRGVYVIK 514
Query: 495 SVPSPKEITFLQKYYMLFGFEVNDYNSFIEPINSMTVCNYLKCTGTYTIRDIDPMLMEQL 554
+ L + +G++ N + + + NY++ I +++ ++++
Sbjct: 515 KQIKEEYRNILSDFSRKYGYKTNLVK--MPNLRTRESYNYVQTKDCNIIGNLNNEDLQKI 572
Query: 555 KAILESGVRFWHNDGSGNPMLQN 577
+ I +SG+ WH D G+ L N
Sbjct: 573 RTIFDSGITLWHADPVGDYTLNN 595
                                                                                >gi|215339 (M12456) p9 tail protein [Bacteriophage phi-29]
            >gi|224163|prf||1011232C protein p9,tail [Bacteriophage
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phi-291 Length = 335

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Score = 71.0 bits (171), Expect = 2e-11
 Identities = 64/293 (21%), Positives = 123/293 (41%), Gaps = 20/293 (6%)
Query: 292 KDEFKHMIRNEYMTIEFYDWNGNTMLLDAGKISQKTGVKLRTKSIIGYHNEVRVYPVDYN 351
KD+ ++ Y E D+ GN M L I+ +K++ ++G N+V DYN
Sbjct: 57 KDQESKLMMYPYCVTEITDFKGNHMNLKTEYINNSK-LKIQVRGSLGVSNKVAYSVQDYN 115
Query: 352 SAENDRPILAKNKEILIDTGSFLNTNITFNSFAQVPILINNGILGQSQQANRQ--KNAES 409
+ D + N+ S +N N I I N L Q N+ +N +S Sbjct: 116 A---DSALSGGNRLTASLDSSLINNNPN------DIAILNDYLSAYLQGNKNSLENQKS 165
Query: 467 LQPPSVTESEMGNAFQIANSINGLTMKISVPSPKEITFLQKYYMLFGFBVNDYNSFIEPI 526 PP +T+ AF N G+ + + L ++ +G+++N + Sbjct: 224 NIPPQLTKMGGNTAFDYGNGYRGVYVIKKQLKAEYRRSLSSFFHKYGYKINRVKK--PNL 281
Query: 527 NSMTVCNYLKCTGTYTIRDIDPMLMEQLKAILESGVRFWHNDGSGNPMLQNPL 579
                 NY++ + DI+ +++++ I ++G+ WH D GN ++N L
Sbjct: 282 RTRKAFNYVQTKDCFISGDINNNDLQEIRTIFDNGITLWHTDNIGNYSVENEL 334
>gi|1181968|emb|CAA87738.1| (Z47794) tail protein (Bacteriophage
           Length = 230
 Score = 53.9 bits (127), Expect = 3e-06
 Identities = 29/113 (25%), Positives = 54/113 (47%), Gaps = 3/113 (2%)
Query: 1 MRKLTNFKFFYNTPF-TDYQNTIHFNSNKERDDYFLNGRHFKSLDYSKQPYNFIRDRMEI 59
           M++ T + +PF DY N I+F + + +D+F + Y
Sbjct: 1 MQESTKIWLYAKSPFKNDYANVINFETRESMEDFFTKKNPHIEIVYEYDKFQYTQRNGSI 60
Query: 60 NVDMQWHDAQGINYMTFLSDFEDRRYYAFVNQIEYVNDVVVKIYFVIDTIMTY 112
'V + + + YM F+++ R YYAFV + Y+N+ +I + +D TY
Sbjct: 61 VVSGRVEKYENVTYMRFINN--GRTYYAFVFDVLYINEDATRIIYEVDVWNTY 111
>gi|1181970|emb|CAA87740.1| (Z47794) tail protein [Bacteriophage
           CP-1]
           Length = 586
 Score = 42.2 bits (97), Expect = 0.010
 Identities = 79/381 (20%), Positives = 139/381 (35%), Gaps = 92/381 (24%)
Query: 277 LSLSFSNLQEMMLSK--KDEFK---HMIRNEYMTIEFYDWNGNTMLLDAG----KISQKT 327
           L +++ +QE + S KD+ + ++ +E+ IE YD GN+ +
Sbjct: 187 LKIAYDQIQEGLRSYMGKDDLEIEVQLLNSEFTEIELYDIYGNSYVYQPQYLPRTIDEAH 246
Query: 328 GVKLRTKSIIGYHNEVRVYPVDYNSAEN----DRPIL----- 360
                  +G N+V + ++YN+A N D+ IL
Sbjct: 247 KYKVIVSGSLGDSNQVHINFLEYNNANNVSYADKNILDSLESGDWAEHNPEHFKYGLNDV 306
Query: 361 -AKNKEILIDT-GSFLNTNITFNSFAQVPILINNGILGQSQQANRQKNAESQLITNRIDN 418
K+ IL D S++ ++ Q+ N +L QS + ++ A + + Sbjct: 307 TGKSVAILNDAEASYIQSHKNQMEHTQLTFKENRDMLKQSVDLSNKQVATANSQASYNAQ 366
Query: 419 VLNGSDPKSRFYDAVSVASNLSPTALFGKF------NEEYNFYKQQQ-- 459
S +++ + S N++ L G F N +YN QQ

Sbjct: 367 FAVDSANINQWTEGASGILNVAGNLLTGNFGGALGGLASGGMKVFNANRDYNDKVVQQGF 426
Query: 460 ------AEYKDLALQPPSVTESEMGNAFQIANSIN 488
                                           A DL QP SV + AFQ N +
Sbjct: 427 TSENNALKSQSNALANMKSKIALDQSIRAYNATMADLQNQPISVQQIGNDLAFQSGNRLT 486
Query: 489 GLTMKISVPSPKEITFLQKYYMLFGFEVNDY-NSFIEPINSMTVCNYLKCTGTY--TIRD 545
+ K+S+ + + + + + + G VN + N + + S DYVKVSLAQKEIMGRANEYIKCYGVLVNWFTNDALSVMRSRKRPNYIKMINVNLGTLR- 545
Query: 546 IDPMLMEQLKAILESGVRFWH 566
            + M ++AI +SGVR W+
Sbjct: 546 ANQSHMNAIQAIFQSGVRIWN 566
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Query= pt|110875 44AHJDORF005 Phage 44AHJD ORF |12643-13890|-1 1
           (415 letters)
>gi|3845203 (AE001399) GAF domain protein (cyclic nt signal
             transduct.) [Plasmodium falciparum]
             Length = 1245
 Score = 52.3 bits (123), Expect = 6e-06
 Identities = 59/246 (23%), Positives = 105/246 (41%), Gaps = 27/246 (10%)
Query: 174 ESIDRNHGNVDYIGFPKMFLLGNAVNFSSPILSNLNIYNLLQKHKMNTSRLYKNIFLEMR 233
+S D N+ N + + N+V FS+ N IY++L N +YK + E+
Sbjct: 854 DSSDNNNNNNNNNNNNNNNNNSVIFST----NEKIYDML-----NRDNIYKKVKKEIF 904
Query: 234 RNDYVNEKRNTRAFNSNDDAMTTGEFEFNEYNLADDNLRNHINQNGDFFYIKTDDKY1-- 291
D + + + + + N + M + N N ++N+ N + N NGD Y KY
Sbjct: 905 EGDSIKTMENKPULTNKNYMNNDNIDNNNNNNNNNNNNNNNDDNIYNDDLKKYYLN 964
Query: 292 KVMYNVTTFMTNIIVVPYTKQYEFCTKIR-DIDNHVTYLRDDMFYKENMERYYYNPSNLH 350
++N ++ + + + K E K+ I + L +F+K NM + + L+
Sbjct: 965 TSIFNKDLYVKHFVDIIMNKSLEEIIKMNVYISERINSL---LFHKGNM----LNDVTKLY 1018
Query: 351 FDNAYSKNYVVDNDRYLYLDMNKIIKFHIKNEMKKNMSEFERKEKIYEDN----YIENTK 406
                                     KIF + BK +M F+ +KIY+ N
                NAY +
Sbjct: 1019 MSNAYGEKCFFFN-----FPQIKEIIFVNEYEKKMDMKYFKMLKKIYKYNLNKIFSNNYK 1073
Query: 407 KYLMKQ 412
               +++K+
Sbjct: 1074 PFIIKK 1079
>gi|3758843|emb|CAB11128.1| (Z98551) predicted using hexExon;
             MAL3P6.23 (PFC0820w), Hypothetical protein, len: 4982 aa
              [Plasmodium falciparum]
             Length = 4981
 Score = 49.2 bits (115), Expect = 5e-05
 Identities = 67/287 (23%), Positives = 110/287 (37%), Gaps = 60/287 (20%)
Query: 127 ITDLNSATDLKYHSNFLKHYPIIIYDEFLALEDDYLIDEWDKLKT----IYESIDRNHGN 182
I D+N + D+ + +++ I YD +++DK++ IY +ID++ N
Sbjct: 3619 IMDINKSKDISKNMEIVQN---IEYD------NKYDKIRNDMDAIYMAIDKDMDN 3664
Query: 183 VDYIGFPKMFLLGNAVNFSSPILSNLNIYNL----LQKHKMNTSRLYKNIFLEMRRNDYV 238
+ I + F L N S +N YNL ++ K N R Y N F +D
Sbjct: 3665 IGIINCMRYFNLYKNYNNLSNECNNRE-YNLNELYMEDIKRNMKR-YDNNFNINHYDDNN 3722
Query: 239 NEKRNTRAFNSNDDAMTTGEFEFNEYNLADDNLRNHINQNGDFFYIKTDDKYIKVMYNVT 298
                                      N N ++N N+ N NG F+ D
              N N N+N++
Query: 299 TFMTNIIVVPYTKQYEFCTKIRDIDNHVTYLRDDMFYKENMERYYYNPSNLHFDNAYSKN 358
                          K FCTK
                                                  ++F +N+E N N N Y+ N
Sbjct: 3772 ------KDLFFCTK------KNIFPCKNIETVCKNEYNKKIYNNYTCN 3807
Query: 359 YVVDNDRYLYLDMNKIIKFHIKNEMKKNMSEFERKEK-IYEDNYIEN 404
                           + ++IK + + N E+ + EK +Y + EN
Sbjct: 3808 ISVNNTLNCLNIIKELIKLNNNKKKILNYYEYHKVEKLLYYRHSFEN 3854
 Score = 35.6 bits (80), Expect = 0.70
 Identities = 62/290 (21%), Positives = 121/290 (41%), Gaps = 65/290 (22%)
Query: 2 VKQNRLDMVRDYQNAVN--HVRKKIPDKYNQIELVDELMNDDIDYYISISNRSDGKSFNY 59
+K+N ++ +N +N +V++ DK N I D++I+ SN + +SF
Sbjct: 4445 IKRNNINKSNIKRNNINKSNVKRSNTDKSNVIS------DFHIT-SNNNITRSFT- 4492
Query: 60 VSFFIYLAIKLDIKFTLLSRHYTLRDAYRDFIEEIIDENPLFKSKRVTFR$ARDYLAIIY 119 -
A D F LS TL +Y +F + + I
Sbjct: 4493 -----ATLTDSIFNTLSE--TLNYSYDNFFSNMDN------IKI 4523
Query: 120 QDKEIGVITDLNSATDLKYHSNFLKHYPIIIYDEFL----ALEDDYLIDEWDKLKTIYE 174
+ EI ITD++ +YH N+LK + +E++ + +D + DE ++T+ E
Sbjct: 4524 KKNEINNITDVDYGNKKEYHENYLKVKONKVNEEYIEETFKSDKDCSIKDEACTIRTLSE 4583
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Query: 175 S--IDRNHGNVDYIGFPKMFLLGNAVNFSSPILSNLNIYNLLQKHKMN--TSRLYKNIFL 230
S I N N+D + + + S P N++ N ++K+ +N R+ KN
Sbjct: 4584 SCNISENISNID------MDDEDHISFPNGRNVHDNNYMKKNHVNYDKMRVGKNKIP 4634
Query: 231 EMRRNDYVNEKRNTRAFNSNDDAMTTGEFEFNEYNLADDNLRNHINQNGD 280
D + +++ + +D M++ ++ E ++ + L + NG+
Sbjct: 4635 SFTHFDKILDEKKK----SDKDMSSSKWLEREEHIKEIKLEKNEYMNGN 4680
 Score = 34.0 bits (76), Expect = 2.0
 Identities = 47/211 (22%), Positives = 84/211 (39%), Gaps = 32/211 (15%)
Query: 210 IYNLLQKHKMNTSRLYKNIFLEMRRNDYVNEKRNTRAFNSNDDAMTTGEFEFNEYNLADD 269
Query: 270 NLRNHINQNGDFFYIKTD---DKYIKVMYNVTTFMTNIIVVPYTKQYBFCTKIRDIDNHV 326 N+ N + D + D+ K MY + V B K D+ N+ Sbjct: 965 NVTNEHGNHSDSYPYGNSLNLDRKPKNMYE-DIYKEKGFVKSDCSNIEI--KKNDMINND 1021
Query: 327 TYLRDDMFYKENMERYYYNPSNLHFDNAYSKNYVVDNDRYLYLDMNKII----KFHIKNE 382
Y +++ FY+++ Y+ + YV++ +YL +N ++ F +KN+
Sbjct: 1022 VYKKNE-FYEDSRINMIYDEDEIKTWFLIPHKYVIN---IIYLFLNILLTDESNFKLKNK 1077
Query: 383 MKKNMSEFERKEKIYEDN-----YIENTKKY 408
E K IYEDN ++N KKY
Sbjct: 1078 KYGYFVNEETKGTIYEDNNGLQEILKNGKKY 1108
 Score = 33.6 bits (75), Expect = 2.7
 Identities = 42/198 (21%), Positives = 77/198 (38%), Gaps = 42/198 (21%)
Query: 222 SRLYKNIFLEMR---RNDYVNEKRNTRAF-----NSNDDAMTTGEFEFNEYNLA 267
S LY I++ + +N + K+NT + N+++D TT B + +
Sbjct: 411 SVLYSIIYMNKKYKKNFIITNKKNTNVYFENDVIQLSVENTSEDTFTTNTRESSLNSGM 470
Query: 268 DDNLRNHINQNGDFFYIKTDDKYIKVMYNVTTFMTNIIVVPYTKQYEFCTKIRDIDNHVT 327
+++R +N D +DDK ++Y N YTK E
Sbjct: 471 MNDMRYSVNNYADEKVYHSDDKSDHLIYKHVHDEKNKYDEMYTKTKE------ 517
Query: 328 YLRDDMFYKENMERYYYNPSNLHFDNAYSKNYVVDNDRYLYLDMNKIIKFHIKNEMKKNM 387
+++ YK N+ + N K LD+ K I H+KN+ + N
Sbjct: 518 --NENIIYKSNIVDKKTCDISSEMVNGKDK------LDVEKYIGSHVKND-ENNK 563
Query: 388 SEFERK-EKIYEDNYIEN 404
              + ++K + + + YI+N
Sbjct: 564 EKLKKKIDNVNKKEYIDN 581
>gi|3845297 (AE001421) hypothetical protein [Plasmodium falciparum]
              Length = 2380
 Score = 48.0 bits (112), Expect = 1e-04
 Identities = 87/390 (22%), Positives = 160/390 (40%), Gaps = 65/390 (16%)
Query: 20 VRKKIPDKYNQIELVDELMNDDIDYYISISNRSDGKSFNYVSFF-----IYLAIKLDIKF 74
+++K +K ++ + +N D + ++ R K+ NY++ +YL I DI
Sbjct: 1049 LQRKNMNKCSKNRNRNRYINKDSNIHLMNLIRIKFKNLNYMNMNSFEIBLYLKINNDIFL 1108
             TLLSRHYTLRDAYR-----DFIEEIIDEN-PLFKSKRVTFRSARDYLAIIYQDKEIGVI 127
+Y +++ Y + + + EN + +++ + + Y +K+
Query: 75
Sbjct: 1109 QFNKHNYNVQNFYNFSITLINIMSKYYSENFYAYNLEKIVYKFLLNNKNFEYIEKQYSSK 1168
Query: 128 TDLNSATDLKYHSNFLKHYPIIIYDEFLA----LEDDYLIDEWDKLKTIYESIDRNHGNV 183
D+N D+ ++ +K+ II EFL L+ D I + KLKT ++
Sbjct: 1169 EDMNEL-DILVNTYDMKYDKII---EFLKNNGYLKIDRYIYFYPKLKT------DI 1214
Query: 184 DYIGFPKMFLLGNAVNFSSPILSNLNIYNLLQKHKMNTSRLY-----KNIF--LEMRRN 235
                   F ++FL N + L NI +++ K + Y
Sbjct: 1215 ILFFFKEIFLNDNILKIDRKFLKK-NITIMIEVLKEIFFKEYVKRCITKVIFFPVHMKEH 1273
Query: 236 DYVNEKR-----NTRAFNSNDDAMTTGEFEFNEYNLADDNLRNHINQNGDFFYIKTD 287
                             N+ FN+ D + N YN D+ N+ N N +Y K
```

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Sbjct: 1274 DHVMNKNYYNNQYVNNSNMFNTRGDHNNNNQTNDNHYNHHYDDTHNNNNNNSKYY-KNK 1332
Query: 288 DKYIKVMYNVTTFMTNIIV---VPYTKQYEFCTKIRDIDNHVTYLRDDMFYKEN----ME 340
                        +++ + V K + K I + Y+ ++ N
             +K K+MY
Sbjct: 1333 NKN-KIMYEKERKSSSLFISNNVQDVKPIKHYLKYSSIYKNFIYIISEIKNFNNKITKIN 1391
Query: 341 RY-YYNPSNLHFDNAYSKNYVVDNDRYLYL 369
            RY YYN NL+ D+ ND YL+L
Sbjct: 1392 RYNYYNYMNLNIDDL-----NDAYLFL 1413
 Score = 32.5 bits (72), Expect = 6.0
 Identities = 46/183 (25%), Positives = 73/183 (39%), Gaps = 26/183 (14%)
Query: 225 YKNIFLEMRRNDYVNEKRNTRAFNSNDDAMTTGEFEFNEYNLADDNLRNHINQNGDFFYI 284
            +KNI ++ ++N + NSN + + N N+ +N N IN +
Sbjct: 27 HKNINKNIKNKKFINIDNSNNCNNSNSNNSNSNNNNNNNNNNIVRNN-NNFINADKKKNVI 85
Query: 285 KTDDKYIKVMYNVTTFMTNIIVVPYTKQYEFCTKIRDIDNHVTYLRDDMFYKENMERYYY 344
+D IK V NI Y ++ + D+ N+ + + KE ER
Sbjct: 86 LNEDDDIKNKELVDESFVNIFF--YENYFKNLFNLNDVSNNKVI--NIIEQKEGDER--- 138
Query: 345 NPSNLHFDNAYSKNYVVDNDRYLYLDMNKIIKFHIKNEMKKNMSEFERKEKIYEDNYIEN 404
N N N +KN V DN +NK IKN +N++E Y N++ +
Sbjct: 139 NADN----NLKNKNIVRDN------INK----IKN--TRNVNEILIYNNKYIINFLND 180
Query: 405 TKK 407
           тк
Sbjct: 181 TTK 183
>qi|4493936|emb|CAB38972.1| (AL034556) predicted using hexExon;
           MAL3P5.6 (PFC0600w), Hypothetical protein, len: 250 aa
           [Plasmodium falciparum]
           Length = 249
 Score = 47.3 bits (110), Expect = 2e-04
 Identities = 53/215 (24%), Positives = 87/215 (39%), Gaps = 30/215 (13%)
Query: 209 NIYNLLQKHKMNTSRLYKNIFLEMRRNDYVNEKRNTRAFNSNDDAMTTGEFEF--NEYNL 266
NIYN L++ YKN N ++ +N N+N EFE N YN
Sbjct: 13 NIYNKLBEK-----YKNFLKLKNMNSHMGASQNMNV-NNNYTMNELEEFEKINNNYNN 64
Query: 267 ADDNLRNHINQNGDFFYIKTD-----DKYIKVMYNVTTFMTNIIVVPYTKQYEFCTKIRD 321
            ++N+ N+IN D+ IK +K ++ YN
Sbjct: 65 NNNNINNNINNYYDYMNIKVSQSVQHNKRLQDFYNNKNSFQHYIKKLKTCRFDADDIRNL 124
Query: 322 IDNHVTYLRDDMFYK-----ENMERYYYNPSNLHFDNAYSKNYVVDNDRYLYLDMNKIIK 376
++ + Y RD+ K EN + N + N + S NY DN + LY +N++ K
Sbjct: 125 LEKRLAYERDNTLIKNIQEENKKGIGINGNFGSESNSSSSNY--DNNYLLYRKINRLNK 182
Query: 377 FHIKNEMKKNMSEFERKEKIYEDNYIENTKKYLMK 411
                             KI

    KKY++K

Sbjct: 183 TNTNKSKNRSRKRKRINSKI------DKKYIIK 209
>gi|3845165 (AE001390) hypothetical protein [Plasmodium falciparum]
           Length = 1247
Score = 45.7 bits (106), Expect = 6e-04
Identities = 52/239 (21%), Positives = 94/239 (38%), Gaps = 38/239 (15%)
Query: 206 SNLNIYNLLQKHKMNTSRLYKNIFLEMRRNDYVNEKRNTRAFNSNDDAMTTGEFEFNEYN 265
           +N N +N ++K K R I +N + +N ++N+D
Sbjct: 474 NNTNKWNEIKKRKKKFKREKNKIINNSFQNQEAEDDKNNNNNDNNNDNHNDNNNENNNEN 533
Query: 266 LADDNLRNHINQNGDFFYI-KTDDKYIK----VMYNVTTFMTNIIVVPYTKQYEFCTKIR 320
             D+N N+ + N D I D+ Y
                                           +YN T ++ YTK + +
Sbjct: 534 NNDNNNENNNDINNIHNNDNNYYNNDNINLYNEMTKKKCMLDNSYTKYFFYIFTL- 592
Query: 321 DIDNHVTYLRDDMFYKENME------RYYYN-------PSNLHFDNAYS 356
                                   ++YYN
              + + ++ + FY++N +
Sbjct: 593 ---DMLPSIKFETFYEKNTDHKNFNENYKFYYNTDDDTDIINAIKKKNVKNKKKNGNIVI 649
Query: 357 KNYVVDNDRYLYLDMNKIIKFHIKNEMKKNMSEFER----KEKIYEDNYIENTKKYLMK 411
           KNY+ \quad N+ \quad Y \quad YL+ \quad N+ \qquad + \quad I \quad + \quad K \qquad +E \qquad \quad K+ \quad I+ \quad ++Y \quad E \qquad K \qquad K
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Sbjct: 650 KNYINHNE-YSYLEYNENKNYEINKKEKLLTENYEYDMYIKDNIHYNDYSEGDGKQTKK 707
 Score = 41.0 bits (94), Expect = 0.016
 Identities = 58/245 (23%), Positives = 96/245 (38%), Gaps = 43/245 (17%)
Query: 207 NLNIYNLLQKHKMNTSRLYKNIFLEMRRNDYVNEKRNTRAFNSNDDAMTTGEFEFNEYNL 266
            N+N+YN + K K Y F + D + +
                                                          + N D
Sbjct: 564 NINLYNEMTKKKCMLDNSYTKYFFYIFTLDMLPSIKFETFYEKNTDHKNFNENYKFYYNT 623
Query: 267 ADD-------NLRNHINQNGDFF---YIKTDDKYIKVMYNVT-TFMTNIIVVPYTKQ 312
DD N++N +NG+ YI ++ Y + YN + N T+
Sbjct: 624 DDDTDIINAIKKKNVKNK-KKNGNIVIKNYINHNE-YSYLEYNENKNYEINKKEKLLTEN 681
Query: 313 YEFCTKIRDIDNHVTYLRDDMFYKENMERYYYNPSNLHFDNAYSK-----NYV--VD 362
YE+ I+D ++ Y D + + YN +N +N Y K +Y+ VD
Sbjct: 682 YEYDMYIKDNIHYNDYSEGDGKQTKXASSFLYNNNN---NNKYKKEDNKTQIISYMDHVD 738
Query: 363 NDR--------YLYLDMNKIIKFHIK-NEM----KKNMSEFERKEKIYEDNYIENTKKY 408
N+ Y + +++ F +K N+M K+ F +E I + +EN K+
Sbjct: 739 NENGVKGLKKRNLFYNNSDQLYNFDVKDNDMIKYEKRQSKNFVEEEFINGNRKMENEDKH 798
Query: 409 LMKQY 413
             LKY
Sbjct: 799 LKKHY 803
Query= pt|110877 44AHJDORF007 Phage 44AHJD ORF |2044-3027|1 1
           (327 letters)
 >gi|1181960|emb|CAA87731.1| (Z47794) connector protein
             [Bacteriophage CP-1]
             Length = 337
  Score = 45.7 bits (106), Expect = 5e-04
  Identities = 44/184 (23%), Positives = 84/184 (44%), Gaps = 13/184 (7%)
 Query: 127 QIHKLYDNCMSGNFVVMQNKPIQYNSDIEIIEHYTDELAEVALSRFSLIMQAKFSK--IF 184
 ++HK ++ +V+N Y I +E + ++LA++ L+ L A+ + IF
Sbjct: 125 ELHKONPDKIKRPCIVIPNNNF-YEPYIGYLELFCEKLADIELT-IQLNRNAQITPYFIF 182
 Query: 185 KSEINDESINQLVSEIYNGAPFVKMSPMFNAD------DDIIDLTSNSVIPALTEMKR 236
                  N S+ + ++I N P V ++ + D D I +
 Sbjct: 183 ADNTNVLSMKNIFNKIANFEPVVYLNKQKDQDGQDSFKQLSDYIQVFRTDAPFLLDKLHD 242
 Query: 237 EYQNKISELSNYLGINSLAVDKESGVSDEEAKSNRGFTTSNSNIYLKGREP-ITFLSKRY 295
 E +++L ++GIN+ DK+ + EA SN G ++N + K R + ++K Y
Sbjct: 243 EKLRVMNQLLTFIGINNNPSDKKERLVVSEAISNNGVISANIEVGWKSRRKFVELINKCY 302
 Query: 296 GLDI 299
             GL+I
 Sbjct: 303 GLEI 306
 >gi|1429239|emb|CAA67658| (X99260) upper collar protein
              [Bacteriophage B103]
              Length = 308
  Score = 44.9 bits (104), Expect = 8e-04
  Identities = 40/159 (25%), Positives = 73/159 (45%), Gaps = 11/159 (6%)
 Query: 150 YNSDIEI----IEHYTDELAEVA-LSRFSLIMQAKFSKIFKSEINDESINQLVSEIYNG 203
 YN+D++ +E + +LAE+ + + Q I ++ N S+ + ++
Sbjct: 121 YNNDLKCSTLPALEMFAQDLAELKEIIAVNQNAQKTPVLIAANDNNQLSLKNIYNQYEGN 180
 Query: 204 APFVKMSPMFNADD-DIIDLTSNSVIPALTEMKREYQNKISELSNYLGINSLAVDKESGV 262
 AP ++ + D+ + + V+ L K N E+ YLGI + ++K+ +
Sbjct: 181 APVIFVHESLDLDNLKVFKTDAPYVVDKLNAQKNAVWN---EVMTYLGIKNANLEKKERM 237
 Query: 263 SDEEAKSNRGFTTSNSNIYLKGR-EPITFLSKRYGLDIK 300
                 E SN
                            S+ NIYLK R E +S+ YGL++K
  Sbjct: 238 VTSEVDSNDEQIESSGNIYLKARQEACNKISELYGLNLK 276
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>gi|137915|sp|P07535|VG10\_BPPZA UPPER COLLAR PROTEIN (CONNECTOR PROTEIN) (LATE PROTEIN GP10) >gi|75851|pir||WMBP10 gene

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10 protein - phage PZA >gi|216059 (M11813) upper collar
            protein [Bacteriophage PZA]
            Length = 309
 Score = 43.8 bits (101), Expect = 0.002
 Identities = 38/160 (23%), Positives = 75/160 (46%), Gaps = 13/160 (8%)
Query: 150 YNSDIEI-----IEHYTDELAEVALSRFSLIMQAKFSKIF--KSEINDESINQLVSEIYN 202
YN+D+ +E + ELAE+ S+ A+ + + + N S+ Q+ ++
Sbjct: 122 YNNDMSFPTTPTLELFAAELAELK-EIISVNQNAQKTPVLIRANDNNQLSLKQVYNQYEG 180
Query: 203 GAPFVKMSPMFNADD-DIIDLTSNSVIPALTEMKREYQNKISELSNYLGINSLAVDKESG 261
AP + ++D ++ + V+ L K N E+ +LGI + ++K+
Sbjct: 181 NAPVIFAHEALDSDSIEVFKTDAPYVVDKLNAQKNAVWN---EMMTFLGIKNANLEKKER 237
Query: 262 VSDEEAKSNRGFTTSNSNIYLKGR-EPITFLSKRYGLDIK 300
            + +R SN S+ ++LK R E
                                           +++ YGLD+K
Sbjct: 238 MVTDEVSSNDEQIESSGTVFLKSREEACEKINELYGLDVK 277
>gi|137914|sp|F04332|VG10_BPPH2 UPPER COLLAR PROTEIN (CONNECTOR
PROTEIN) (LATE PROTEIN GP10) >gi|75852|pir||WMBPC9 gene
10 protein - phage phi-29 >gi|215328 (M14782) upper
            collar protein (Bacteriophage phi-29) >gi|215340 (M12456) pl0 connector protein (Bacteriophage phi-29)
            >gi|224161|prf||1011232A protein pl0, connector
            [Bacteriophage phi-29] >gi|225365|prf||1301270E gene 10
            (Bacteriophage phi-29)
            Length = 309
 Score = 41.4 bits (95), Expect = 0.009
 Identities = 37/160 (23%), Positives = 75/160 (46%), Gaps = 13/160 (8%)
Query: 150 YNSDIEI-----IEHYTDELAEVALSRFSLIMQAKFSKIF--KSEINDESINQLVSEIYN 202
YN+D+ +E + ELAE+ S+ A+ + + + N S+ Q+ ++
Sbjct: 122 YNNDMAFPTTPTLELFAAELAELK-EIISVNQNAQKTPVLIRANDNNQLSLKQVYNQYEG 180
Query: 203 GAPFVKMSPMFNADD-DIIDLTSNSVIPALTEMKREYQNKISELSNYLGINSLAVDKESG 261
AP + ++D ++ + V+ L K N E+ +LGI + ++K+
Sbjct: 181 NAPVIFAHEALDSDSIEVFKTDAPYVVDKLNAQKNAVWN---EMMTFLGIKNANLEKKER 237
Query: 262 VSDEEAKSNRGFTTSNSNIYLKGR-EPITFLSKRYGLDIK 300 + +E SN S+ ++LK RE +++ YGL++K Sbjct: 238 MVTDEVSSNDEQIESSGTVFLKSREEACEKINELYGLNVK 277
Query= pt|110878 44AHJDORF008 Phage 44AHJD ORF |3020-3775|2 1
          (251 letters)
>gi|4982468|gb|AAD30963.2| (AF118151) SNF1/AMP-activated kinase
            [Dictyostelium discoideum]
 Score = 52.3 bits (123), Expect = 3e-06
 Identities = 28/118 (23%), Positives = 56/118 (46%), Gaps = 5/118 (4%)
Query: 177 TTLRFADNNTIDNGKTVNKSSNESNQNAKRNQNQKGNAKGTQFTKQYLID-NIDKAYD 233 +NN I+N N N ++N +N N N N N N+ + T+ + I N++ +Y+
Score = 37.5 bits (85), Expect = 0.094
 Identities = 17/111 (15%), Positives = 45/111 (40%)
Query: 130 HNEDTTSNTDETSNQNATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNTIDN 189
            +N + +N + +N N + +N++ ++ + P + ++++
Query: 190 GKTVNKSSNESNQNAKRNQNQKGNAKGTQFTKQYLIDNIDKAYDLRKKILN 240
               N +N +N N N N
                                                ID+++ + + + N
Sbjct: 516 NNNTHNDNNNNNNNNNNNNNNNNNNNNNNNNNNCIDSVNNSLNNENDVNN 566
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Score = 32.8 bits (73), Expect = 2.4
  Identities = 31/140 (22%), Positives = 57/140 (40%), Gaps = 14/140 (10%)
 Query: 166 PQSEVN--IDVDNTTLRFADNNTIDNGKTVNKSS-----NESNQNAKRNQNQKGNAK 215
 + +N DV+N+ + +NN D+G N ++ N N + N GN
Sbjct: 554 VNNSLNNENDVNNSNINNNNNNSDDGSNNNSYEGGGDVLLLSDLNGNNQLGGNDNGNVV 613
 Query: 216 GTQFTKQYLIDNIDKAYDLR 235
Q L++++D D++
Sbjct: 614 NLNNNFQ-LLNSLDLNSDIQ 632
 Score = 31.7 bits (70), Expect = 5.4
 Identities = 25/115 (21%), Positives = 48/115 (41%), Gaps = 10/115 (8%)
Query: 130 HNEDTTSNTDETSNQNATSLDNST---GMTAN-RNAYVSLPQSEVNIDVDNTTLRFADNN 185
+N + +N + +N N +S+ T ++ N N+Y S S N + N+ +N
Sbjct: 462 NNNNNNNNNNNNNNNNNSISGGTEVFSISPNLNNSYNS--NSSGNSNSNNNNNNNNT 519
Query: 186 TIDNGKTVNKSSNESNQNAKRNQNQKGNAKGTQFTKQYLIDNIDKAYDLRKKILN 240
Score = 31.7 bits (70), Expect = 5.4
 Identities = 15/104 (14%), Positives = 43/104 (40%)
Query: 110 NVVYSSSEVEKYLQSQGFTEHNEDTTSNTDETSNQNATSLDNSTGMTANRNAYVSLPQSE 169
Query: 170 VNIDVDNTTLRFADNNTIDNGKTVNKSSNESNQNAKRNQNQKGN 213
+N ++ ++ ++ +N N +++ +N N N N N Sbjct: 494 LNNSYNSNSSGNSNGSNSNNNSNNNTNNDNNNNNNNNNNNNNNN 537
 Score = 30.9 bits (68), Expect = 9.2 Identities = 16/84 (19%), Positives = 34/84 (40%)
Query: 130 HNEDTTSNTDETSNQNATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNTIDN 189
          +N + +N + +N N + +N+ +
                                         S+ + N N++
Query: 190 GKTVNKSSNESNQNAKRNQNQKGN 213
                + N +N N N N
Sbjct: 515 SNNNTNNDNNNNNNNNNNNNNNNN 538
>gi|1730077|sp|P18160|KYK1_DICDI NON-RECEPTOR TYROSINE KINASE SPORE
          LYSIS A (TYROSINE-PROTEIN KINASE 1) >gi|974334 (U32174)
          non-receptor tyrosine kinase [Dictyostelium discoideum]
          Length = 1584
 Score = 46.5 bits (108), Expect = 2e-04
 Identities = 29/106 (27%), Positives = 48/106 (44%), Gaps = 4/106 (3%)
Query: 186 TIDNGKTVNKSSNESNQNAKRNQNQKGNAKGTQFTKQYLIDNIDKA 231
+N N +SN +N N N N N TK+ I + D++
Sbjct: 502 NNNNNNNSNSNSNNNNINNNNNNNNNNNNNNNIYLTKKPSIGSTDES 547
                                                                          Score = 34.0 bits (76), Expect = 1.1
 Identities = 20/117 (17%), Positives = 46/117 (39%)
Query: 87 NRQTVEAFGMQVITVCITHEDYLNVVYSSSEVEKYLQSQGFTEHNEDTTSNTDETSNQNA 146
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G IT T + + ++ + + +N + +N +

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Query: 147 TSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNTIDNGKTVNKSSNESNQN 203
                                               N ++N +N N
          + ++++ T N N + + N + +N
                                       N+ +N
Score = 33.2 bits (74), Expect = 1.8
 Identities = 18/88 (20%), Positives = 35/88 (39%)
Query: 190 GKTVNKSSNESNQNAKRNQNQKGNAKGT 217
            N ++N +N N+ + N T
Score = 32.5 bits (72), Expect = 3.1
 Identities = 18/94 (19%), Positives = 37/94 (39%)
Query: 120 KYLQSQGFTEHNEDTTSNTDETSNQNATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTL 179
         K + S N + +N++ +N N ++ + +T S
Sbjct: 392 KNVNSTSILVPNGNNNNNNNNNNNNNNNNNNIIGNGKITTTTTSTSPSSINNNEDISSNNN 451
Query: 180 RFADNNTIDNGKTVNKSSNESNQNAKRNQNQKGN 213
           +NN +N N ++N N + +
Score = 32.5 bits (72), Expect = 3.1
 Identities = 24/110 (21%), Positives = 44/110 (39%), Gaps = 10/110 (9%)
Query: 138 TDETSNQNATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNTIDNGK----- 191
Query: 192 ----TVNKSSNESNQNAKRNQNQKGNAKGTQFTKQYLIDNIDKAYDLRKK 237
T N +SN +N N N N N N+ +N + L KK
>gi|3758855|emb|CAB11140.1| (Z98551) predicted using hexExon;
         MAL3P6.11 (PFC0760c), Hypothetical protein, len: 3395 aa
         (Plasmodium falciparum)
         Length = 3394
 Score = 46.5 bits (108), Expect = 2e-04
 Identities = 52/202 (25%), Positives = 96/202 (46%), Gaps = 32/202 (15%)
Query: 21 FNEFVNDNKLTFYDDEFQFMQKMLKFD-KDVLAIVNEKVFKGFSLKDELSDL--LFKKSF 77
F ++ ++ K T D+ M+K K D DV + NEK++ L ++L+ + + KK
Sbjct: 665 FEKYCSNIKNTLIRDD---MKKFRKPDISDVHILHNEKIYLEKLLNYIKDIEKKLD 721
Query: 78 TIHFLORBINRQTVEAFGMQV-----ITVCITHEDYLNVVYSSSEVEKYLQSQGFTEHNE 132
+ H + IN+ + + + + QV I V + DY + S + + K + + N
Sbjct: 722 ELHGV---INKNKEDIYILQVEKQTLIKVISSVYDYTKME-SENHIFKMNTTWNKMLNNV 777
Query: 133 DTTSNTDETSNQNATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNTIDNGKT 192
          +SN D +NQN +++N+ + N+N N +++N + N +N
Sbjct: 778 HMSSNKDY-NNQNNQNIENNQNIENNQN------NQNIEN-----NQNIENNQNN 820
Query: 193 VNKSSNESNQNAKRNQNQKGNA 214
         N +N++NQN + NQN + NA
Sbjct: 821 QNNQNNQNNQNNQNNQNNQNNA 842
 Score = 33.6 bits (75), Expect = 1.4
                                                                Identities = 46/221 (20%), Positives = 89/221 (39%), Gaps = 37/221 (16%)
Query: 10 DFIKSELIKKGFNEFVNDNKLTFYDDEFQFMQKMLKFDKDVLAIVNEKVFKGFSLKDELS 69
         D +K E K N + +L Y + + M+K K + V K SL
Sbjct: 367 DSLKIEYNKSKTNIQQLNEQLVNYKNFIKEMEKKYK-----QLVVKNNSLFSITH 416
Query: 70 DLLFKKSFT1HFLDREINRQTVEAFGMQVITVCITH---EDYLNVVYSSSEVEKYLQSQG 126
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D + K+ I + R + + + + + + + H + D+L+V+Y + + L +
 Sbjct: 417 DFINLKNSNIIIIRRTSDMKQI----FKMYNLDIEHFNEQDHLSVIY----IYEILYNTN 468
 Query: 127 FTEHNEDTTSNTDETSNQNATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNT 186
               +N D +N D +N N + +N+ N N N + +N +
 Query: 187 IDNGKTVNKSSNESNQNAKRNQNQKGNAKGTQFTKQYLIDN 227
            I+N + N +++ N + N N + N + ++Y I+N
 Sbjct: 513 IENMNSGNHPNSNNLHNYRHNTNDENNLSSLKTSFRYKINN 553
  Score = 32.8 bits (73), Expect = 2.4
  Identities = 28/122 (22%), Positives = 53/122 (42%), Gaps = 2/122 (1%)
 Query: 119 EKYLQSQGFTEHNEDTTSNTDETSNQNATSLDNSTGMTANRNAYVSLPQSEVNID-VDNT 177
Query: 178 TLRFADNNTIDNGKTVNKSSNESNQNAKRNQNQKGNAKGTQFTKQYLIDNIDKAYDLRKK 237
                  +N+ +NG SSN ++ N N N K N +G + + + + YD K
Sbjct: 2898 NNNDNNNDNSNNGFVCELSSNINDFNNILNVN-KDNFQGINKSNNFSTNLSEYNYDAYVK 2956
Query: 238 IL 239
Sbjct: 2957 IV 2958
 Score = 32.5 bits (72), Expect = 3.1
 Identities = 46/249 (18%), Positives = 101/249 (40%), Gaps = 31/249 (12%)
Query: 9 YDFIKSELIKKGFNEFVNDNKLTFYDDEFQFMQKMLKFDKDVLAIVNEKVFKGFSLKDEL 68
Y+++K ++ N N NK E Q++ K+ + + + + E K L++
Sbjct: 2150 YNYVK---VQNATNREDNKNK-----ERNLSQEIYKYINENIDLTSELEKKNDMLENYK 2200
Query: 69 SDL-----LPKKSFTIHFLDREINRQTVEAFGMQVITVCITHEDYLNVVYSSSEVEKYL 122
++L ++K + I L + M+ ++ N + E+ + L
Sbjct: 2201 NELKEKNEEIYKLNNDIDMLSNNCKKLKESIMMMEKYKIIMN----NNIQEKDEIIENL 2255
Query: 123 QSQGFTEHNEDTTSNTDETSNQNATSLDNSTGMTAN----RNAYVSLPQSE----VNIDV 174 +++ + +D +N + ++S M+ + N + +L +S N+D+ Sbjet: 2256 KNK-YNNKLDDLINNYSVVDKSIVSCFEDSNIMSPSCNDILNVFNNLSKSNKKVCTNMDI 2314
Query: 175 DNTTLRFADNNTIDNGKTVNKSSNESNQNAKRNQNQKGNAKGTQFTKQYLIDNIDKAYDL 234 N + ++I+N +N +N +N +N N N N N K YL++N+ D
Sbjct: 2315 CNENMDSI--SSINNVNNINNVNNINNVNNINNVNNINNVKNIVDINNYLVNNLQLNKDN 2372
Query: 235 RKKILNEFD 243
               I+ +F+
Sbjct: 2373 DNIIIIKFN 2381
 Score = 32.1 bits (71), Expect = 4.1
 Identities = 20/103 (19%), Positives = 48/103 (46%), Gaps = 2/103 (1%)
Query: 115 SSEVEKYLQSQGFTEHNEDTTSNTDETSNQN--ATSLDNSTGMTANRNAYVSLPQSEVNI 172
            +++ EKY
                         EH + ND +N+N
Sbjct: 3264 NNDEEKYSCHDDKNEHTNNDLLNIDHDNNKNNITDELYSTYNVSVSHNKDPSNKENEIQN 3323
Query: 173 DVDNTTLRFADNNTIDNGKTVNKSSNESNQNAKRNQNQKGNAK 215
                     D N ++ N ++E+++N + ++N + + K
Sbjct: 3324 LISIDSSNENDENDENDENDENDENDENDENDENDEN 3366
 Score = 30.9 bits (68), Expect = 9.2
 Identities = 27/118 (22%), Positives = 53/118 (44%), Gaps = 15/118 (12%)
Query: 104 THEDYLNVVYSSSEV----EKYLQSQGFTEHNEDTTSNTDETSNQNATSLDNSTGMTANR 159 T+ D LN+ + ++++ E Y HN+D ++ +E QN S+D+S N
T+ D LN+ + +++ E Y HN+D ++ +E QN S+D+S N
Sbjct: 3280 TNNDLLNIDHDNNKNNITDELYSTYNVSVSHNKDPSNKENEI--QNLISIDSSNENDEND 3337
Query: 160 NAYVSLPQSEVNIDVDNTTLRFADNNTIDNGKTVNKSSNESNQNAKRNQNQKGNAKGT 217
                   +++ N + D D N ++ N +E+++N + ++N N +GT
```

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>gi|585795|sp|P21538|REB1_YEAST DNA-BINDING PROTEIN REB1 (QBP)
            >gi|626139|pir||S45907 DNA-binding protein REB1 - yeast
            (Saccharomyces cerevisiae) >gi|536280 emb|CAA84992|
            (235918) ORF YBR049c [Saccharomyces cerevisiae]
            >gi|559944|emb|CAA86391| (Z46260) REB1 DNA-binding
           protein [Saccharomyces cerevisiae]
            Length = 810
 Score = 45.7 bits (106), Expect = 3e-04
 Identities = 34/158 (21%), Positives = 72/158 (45%), Gaps = 14/158 (8%)
Query: 83 DREINROTVEAFGMQVITVCITHEDYLNVVYSSSEVEKYLQSQGFTEHNEDTTSNTDETS 142
           D+ N+++VE ++ + V + H+++ +++
           DKNANQESVEEAVLKYVGVGLDHQNHDPQLHTKDLENKHSKKQNIVESSSDVDVNNNDDS 66
Query: 143 NQNATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNTID---NGKTVNKSSNE 199
N+N + D+S ++A L +E + +VD+ N +D N+ +E
Sbjct: 67 NRNEDNNDDSENISA-----LNANESSSNVDHANSNEQHNAVMDWYLRQTAHNQQDDE 119
Query: 200 SNQNAKRNQNQKGNAKGTQFTKQYLIDNIDKAYDLRKK 237
++N N GN F++ ++ +D D KK
Sbjct: 120 DDEN--NNYTDNGNDSNNHFSQSDIV--VDDDDDKNKK 153
>gi|172372 (M58728) DNA-binding protein [Saccharomyces cerevisiae]
           Length = 809
 Score = 45.7 bits (106), Expect = 3e-04
 Identities = 34/158 (21%), Positives = 72/158 (45%), Gaps = 14/158 (8%)
Query: 83 DREINRQTVEAFGMQVITVCITHEDYLNVVYSSSEVEKYLQSQGFTEHNEDTTSNTDETS 142
           D+ N+++VE ++ + V + H+++ +++
                                                  K+ + Q E + D N ++ S
           DKNANQESVEEAVLKYVGVGLDHQNHDPQLHTKDLENKHSKKQNIVESSNDVDVNNNDDS 66
Query: 143 NQNATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNTID---NGKTVNKSSNE 199
N+N + D+S ++A L +E + +VD+ N +D N+ +E
Sbjct: 67 NRNEDNNDDSENISA-----LNANESSSNVDHANSNEQHNAVMDWYLRQTAHNQQDDE 119
Query: 200 SNQNAKRNQNQKGNAKGTQFTKQYLIDNIDKAYDLRKK 237
++N N GN F++ ++ +D D KK
Sbjct: 120 DDEN--NNNTDNGNDSNNHFSQSDIV--VDDDDDKNKK 153
>gi|2952545 (AF051898) coronin binding protein [Dictyostelium
           discoideuml
           Length = 560
 Score = 44.9 bits (104), Expect = 6e-04
 Identities = 26/83 (31%), Positives = 39/83 (46%), Gaps = 5/83 (6%)
Query: 131 NEDTTSNTDETSNONATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNTIDNG 190
           N + +N +N N+ S +NS +N N+ + P N D DN T +NNT +N
Sbjct: 404 NNNNNNNIINNNNSNSNNNNSNN-NSNNNSNRNSPNHNNNGDNDNNT----NNNTNNNN 458
Query: 191 KTVNKSSNESNQNAKRNQNQKGN 213
              N ++N +N N N N
Sbjct: 459 NNNNNNNNNNNNNNNNNNNNNN 481
Score = 41.4 bits (95), Expect = 0.006
Identities = 22/88 (25%), Positives = 43/88 (48%), Gaps = 6/88 (6%)
Query: 130 HNEDTTSNTDETSNQNATSLDN---STGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNT 186 + ++ +N++ SN N+ + +N + G AN++ + P + +N + DN +NN Sbjct: 337 NRNNSNNNSNNNSNNRNNRNTTNGSNANKS---NSPNNNLNTNNDNKNNNSNNNNN 393
Query: 187 IDNGKTVNKSSNESNQNAKRNQNQKGNA 214
            +N
                    S+N +N N N N
Sbjct: 394 SNNNSNNGNSNNNNNNNIINNNNSNSNS 421
                                                                                 Score = 40.6 bits (93), Expect = 0.011
Identities = 24/101 (23%), Positives = 41/101 (39%), Gaps = 2/101 (1%)
Query: 115 SSEVEKYLQSQGFTEHNEDTTSNTDETSNQNATSLDNSTGMTANRNAYVSLPQSEVNIDV 174
                L + ++N +N ++ N S +N+ N N S + N +
```

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Query: 175 DNTTLRFADN--NTIDNGKTVNKSSNESNQNAKRNQNQKGN 213
           +N + R + N N DN
                              N ++N +N N N N
Sbjct: 430 NNNSNRNSPNHNNNGDNDNNTNNNNNNNNNNNNNNNNNNN 470
 Score = 40.2 bits (92), Expect = 0.014
 Identities = 21/80 (26%), Positives = 39/80 (48%), Gaps = 9/80 (11%)
Query: 130 HNEDTTSNTDETSNQNATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNTIDN 189
                                                N + +N
           +N D +NT+ +N N + +N+
                                    N N
Query: 190 GKTVNKSSNESNQNAKRNQN 209
             + N +SN +N N +N+N
Sbict: 493 SNSNNNNSNSNNNNDNKNEN 512
 Score = 39.5 bits (90), Expect = 0.024
 Identities = 26/111 (23%), Positives = 44/111 (39%), Gaps = 20/111 (18%)
Query: 112 VYSSSEVEKYLQSQ--GFTEHNEDTTSNTDETSNQNATSLDNSTGMTANRNAYVSLPQSE 169
           VY + K+ ++ G +N ++ +N++ SN N ++N
Sbjct: 296 VYCTHHHTKFYETHRNGLLNNNNNSNNNSNSNSNNNNNGINNRNNSNNNSN----- 346
Query: 170 VNIDVDNTTLRFADNNTIDNGKTVNKSS-----NESNQNAKRNQNQKGNA 214
                     ++N I NG NKS+ N +N'N N N N+
Sbjct: 347 ---NNSNNNSNNSNNRNITNGSNANKSNSPNNNLNTNNDNKNNNSNNNNNS 394
 Score = 37.5 bits (85), Expect = 0.094
 Identities = 24/96 (25%), Positives = 41/96 (42%), Gaps = 1/96 (1%)
Query: 124 SQGFTEHNEDTTSNTDETSNQNATSLDNSTGM-TANRNAYVSLPQSEVNIDVDNTTLRFA 182
Query: 183 DNNTIDNGKTVNKSSNESNQNAKRNQNQKGNAKGTQ 218
+NN DN + +SN +N N+ N + K Q
Sbjct: 481 NNNYADNSNNNSSNSNNNNSNSNNNNDNKNENSDNQ 516
 Score = 35.6 bits (80), Expect = 0.36 Identities = 25/99 (25%), Positives = 42/99 (42%), Gaps = 18/99 (18%)
Query: 130 HNEDTTSNTDETSNQNATSLDNST-GMTANRNAYVSLPQSEVNIDVDNTTLRFADNNTID 188
+N + SN + +N N ++ N T G AN++ + P + +N + DN +NN +
Sbjct: 339 NNSNNNSNNNSNNSNNSNNRNITNGSNANKS---NSPNNNLNTNNDNKNNNSNNNNNSN 395
Query: 189 NGKTV------NKSSNESNQNAKRNQNQKGN 213
Sbjct: 396 NNSNNGNSNNNNNNNIINNNNSNSNSNNNSNNNSNNNSN 434
 Score = 35.2 \text{ bits } (79), \text{ Expect = } 0.47
 Identities = 21/94 (22%), Positives = 42/94 (44%), Gaps = 5/94 (5%)
Query: 124 SQGFTEHNEDTTSNTDETSNQNATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFAD 183 + G + ++ +N T+N N + N+ N N+ + N N+ +N + + + Sbjct: 362 TNGSNANKSNSPNNNLNTNNDNKNNNSNN-----NNNSNNNSNNGNSNNNNNNNIINNNN 416
Query: 184 NNTIDNGKTVNKSSNESNQNAKRNQNQKGNAKGT 217
          +N+ N + N S+N SN+N+ + N N T
Sbjct: 417 SNSNSNNNSNNNSNNNSNRNSPNHNNNGDNDNNT 450
Score = 35.2 bits (79), Expect = 0.47
Identities = 29/118 (24%), Positives = 53/118 (44%), Gaps = 12/118 (10%)
Query: 115 SSEVEKYLQS-QGFTEHNEDTTSNTDETSNQNATSLDNSTGMTANRNAYVSLPQSEVNID 173
          SS+ E ++ +GF + + T+N ++N
                                           D S+G + + + V+ P+S +N
Sbjct: 114 SSDSEADIEDDKGFQD--KPITTNNSGSNNPLKNLKDYSSGSSGSSRSGVNQPRSNINNS 171
Query: 174 VDNTTLRFADNNT-----IDNGKTVNKSSNESNQNAKRNQNQKGNAKGTQFTKQ 222
           D + + +N+
                             I + T + NQN +NQNQ N Q +Q
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Sbjct: 172 NDKYKSKSSSNSNSSSSGGSLISSLLTGGNTYQNQNQNQNQNQNQNNNQSQLQQQQQ 229
  Score = 34.4 bits (77), Expect = 0.81
 Identities = 24/94 (25%), Positives = 38/94 (39%), Gaps = 12/94 (12%)
 Query: 131 NEDTTSNTDETSNQNATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNTIDNG 190
             N +T +N + +N N + +N+ N N S N N
 Sbjct: 451 NNNTNNNNNNNNNNNNNNNNNNNNNNNNNNYADNSNNNSSNSN-----NNNSNSNN 504
Query: 191 KTVNKSSNESNQNAKR-----NQNQKGNAKGTQ 218
NK+ N NQ+ R ++NQK + Q
Sbjct: 505 NNDNKNENSDNQSVLRSNEKFTDENQKNGSDDQQ 538
 Score = 33.6 bits (75), Expect = 1.4
 Identities = 22/90 (24%), Positives = 35/90 (38%)
Query: 124 SQGFTEHNEDTTSNTDETSNQNATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFAD 183
                  N SN +++++ N N+ N N + + N + +N
Sbjct: 353 SNNSNNRNITNGSNANKSNSPNNNLNTNNDNKNNNSNNNNNNNNNNNNNNNNNNNNN 412
Query: 184 NNTIDNGKTVNKSSNESNQNAKRNQNQKGN 213
           NN N + N S+N SN N+ RN
Sbjct: 413 NNNNSNSNSNNNSNNNSNRNSPNHNN 442
>gi|535260|emb|CAA82996| (Z30339) STARP antigen [Plasmodium
            reichenowi]
            Length = 655
 Score = 44.5 bits (103), Expect = 7e-04
 Identities = 31/114 (27%), Positives = 47/114 (41%), Gaps = 14/114 (12%)
Query: 128 TEHNEDTTSNTDETSNQNATSLDNSTGMTANRNAYVSLPQSEVN-----IDVDNTTLRF 181
T++N T TD + + +N+T A N + ++ N D +NT +
Sbjct: 433 TDNNNTNTKATDSNNTNTKATDNNNTNTKATDNNNTNTKATDNNNTNTKA 492
Query: 182 ADNNTI-----DNGKTVNKSSNESNQNAKRNQNQKGNAKGT---QFTKQYLIDN 227 DNN DN T K+++ +N N K N N K T T QY+ N Sbjct: 493 TDNNNTNTKATDNNNTNTKATDNNNTNTKATDNNNTNTKATDNNNTNTKATDNNNTNTKATDNNNTNTRYVFAN 546
 Score = 44.5 bits (103), Expect = 7e-04
 Identities = 30/103 (29%), Positives = 44/103 (42%), Gaps = 13/103 (12%)
Query: 128 TEHNEDTTSNTDETSNQNATSLDNS----TGMTANRNAYVSLPQSEVN----IDVDNTTL 179 T++N T TD+++N + + DN+ T T N N S D +NT
Sbjct: 401 TDNNNTDTKATDKSNNTDTKATDNNNNTDTKATDNNNTNTKATDSNNTNTKATDNNNTNT 460
Query: 180 RFADNNTI-----DNGKTVNKSSNESNQNAKRNQNQKGNAKGT 217 + DNN DN T K+++ +N N K N N K T Sbjct: 461 KATDNNNTNTKATDNNNTNTKATDNNNTNTKATDNNNTNTKAT 503
 Score = 42.6 bits (98), Expect = 0.003
 Identities = 27/96 (28%), Positives = 43/96 (44%), Gaps = 10/96 (10%)
Query: 128 TEHNEDTTSNTDETSNQNATSLD-NSTGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNT 186
            T++N +T + + + N N + D N+T A N + ++ N NT + DNN
Sbjct: 422 TDNNNTDTKATDNNNTNTKATDSNNTNTKATDNNNTNTKATDNN----NTNTKATDNNN 477
Query: 187 I----DNGKTVNKSSNESNQNAKRNQNQKGNAKGT 217
                  DN T K+++ +N N K N N K T
Sbjct: 478 TNTKATDNNNTNTKATDNNNTNTKATDNNNTNTKAT 513
 Score = 41.8 bits (96), Expect = 0.005
 Identities = 35/150 (23%), Positives = 59/150 (39%), Gaps = 9/150 (6%)
                                                                                           Query: 85 EINRQTVEAFGMQVITVCITHEDYLNVVYSSSEVEKYLQSQGFTEHNEDTTSNTDETSNQ 144
E N+ ++ G T+ + N + E ++Q T+N TT+ + N
Sbjct: 118 ETNKTNIKLTGNNSTTINTNLTENTNA--TKKLTENVITNQILTGNNNTTTNTSSTEHNN 175
Query: 145 NATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNTIDNGKTVNKSSNESNQNA 204
            N + NSTG T+
                                        NI + N L +N T + T + ++ +N N+
```

Sbjct: 176 NINTNTNSTGNTSTTKKLTE-----NI-ITNOILTGNNNTTTNTSSTEHNNNINTNTNS 228

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Query: 205 KRNQNQKGNAKGTQFTKQYLIDNIDKAYDL 234
             N N N
                          T + DNI+
Sbjct: 229 TDNSNTNTNLTDITTTTKKWTDNINTTONL 258
 Score = 41.8 bits (96), Expect = 0.005
 Identities = 30/101 (29%), Positives = 43/101 (41%), Gaps = 13/101 (12%)
Query: 130 HNEDTTSNTDETSNQNATSLDNS-TGMTANRNAYVSLPQSEVNIDV-----DNTTLRFA 182
            +N DT S ++ ++ AT DN+ T T N N +
Sbjct: 363 NNTDTISTDNDNTDTKATDNDNTDTKATDNNNTDTKATDNNNTDTKATDKSNNTDTKAT 422
Query: 183 DNN-----TIDNGKTVNKSSNESNQNAKRNQNQKGNAKGT 217
DNN DN T K+++ +N N K N N K T
Sbjct: 423 DNNNNTDTKATDNNNTNTKATDSNNTNTKATDNNNTNTKAT 463
 Score = 40.6 bits (93), Expect = 0.011
 Identities = 31/121 (25%), Positives = 47/121 (38%), Gaps = 31/121 (25%)
Query: 128 TEHNEDTTSNTDETSNQNAT-----SLDNSTGMTANRNAYVSLPQSEVN------- 171
TEHN + +NT+ T N + T ++ + +T N N + +E N
Sbjct: 171 TEHNNNINTNTNSTGNTSTTKKLTENIITNQILTGNNNTTTNTSSTEHNNNINTNTNSTD 230
Query: 172 -----IDVDNTTLRFADN------NTIDNGKTVNKSSNESNQNAKRNQNQKGNAKG 216
                    D+ TT ++ DN T N TV+ +N +N N K N N K
Sbjct: 231 NSNTNTNLTDITTTTKKWTDNINTTQNLTTSTNTTTVSTDNNNNNINTKPTDNNNTNIKS 290
Query: 217 T 217
Sbjct: 291 T 291
 Score = 38.3 bits (87), Expect = 0.055
 Identities = 28/98 (28%), Positives = 41/98 (41%), Gaps = 10/98 (10%)
Query: 128 TEHNEDTTSNTDETSNQNATSLDNSTGMTANRNAYVSLPQSEVNIDVD-NTTLRFADNNT 186
           TEHN + +NT+ S N+ + N T +T +
Sbjct: 216 TEHNNNINTNTN--STDNSNTNTNLTDITTTTKKWTDNINTTQNLTTSTNTTTVSTDNNN 273
Query: 187 -----IDNGKTVNKSSNESNQNAKRNQNQKGNAKGT 217
                   DN T KS++ N K N+ + K T
Sbjct: 274 NNINTKPTDNNNTNIKSTDNYNTGTKETDNKNTDIKAT 311
 Score = 37.5 bits (85), Expect = 0.094
 Identities = 31/106 (29%), Positives = 45/106 (42%), Gaps = 18/106 (16%)
Query: 128 TEHNEDTTSNTDETSNQN----ATSLDNSTGMTANRNAYVSLPQSEVN------IDVDN 176 T++N +T +T T N N AT N+T A N + ++ N D +N Sbjct: 390 TDNNNTT--DTKATDNNNTDTKATDKNNTDTKATDNNNTTKATDSNN 447
Query: 177 TTLRFADNN-----TIDNGKTVNKSSNESNQNAKRNQNQKGNAKGT 217
T + DNN DN T K+++ +N N K N N K T
Sbjct: 448 TNTKATDNNNTNTKATDNNNTNTKATDNNNTNTKATDNNNTNTKAT 493
Score = 35.2 bits (79), Expect = 0.47
Identities = 24/109 (22%), Positives = 46/109 (42%), Gaps = 6/109 (5%)
Query: 128 TEHNEDTTSNTDETSNQNATSLDNSTGMTANRNAYVSLPQSEVN-----IDVDNTTLRF 181
           T++N T TD + + +N+T A N + ++ N D +NT +
Sbjct: 473 TDNNNTNTKATDNNNTNTKATDNNNTNTKATDNNNTNTKATDNNNTNTKATDNNNTNTKA 532
Query: 182 ADMNTIDNGKTVNKSSNESNQNAKRNQNQKGNAKGTQFTKQYLIDNIDK 230
                                                                                 - ______
            DNN N + +E+ + K N++ N++ + K + +DK
Sbjct: 533 TDNNNNTNQYVFANNYDETTSDDKLNKDSCDNSEEKENIKSMINAYLDK 581
Score = 34.4 bits (77), Expect = 0.81
Identities = 26/126 (20%), Positives = 46/126 (35%), Gaps = 7/126 (5%)
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Query: 99 ITVCITHEDYLNVVYSSSEVEKYLQSQGFTEHNEDTTSNTDETSNQNATSLDNSTGMTAN 158
            IT T+ + ++ S + V S T +++ +N T N N ++
Sbjct: 318 ITTDNTNTNVISTDNSKTNVISKDNSNTHTISTDNSKTNVISTDNNNTDTISTDNDNTDT 377
Query: 159 RNAYVSLPQSEVNIDVDNTTLRFADNNTID------NGKTVNKSSNESNQNAKRNQNQK 211 + ++ ++ +NT + DNN D N + N +N + K N
Sbict: 378 KATDNDNTDTKATDNNNTDTKATDNNNTDTKATDKSNNTDTKATDNNNTDTKATDNNN 437
Query: 212 GNAKGT 217
            NKT
Sbjct: 438 TNTKAT 443
 Score = 34.4 bits (77), Expect = 0.81
 Identities = 30/100 (30%), Positives = 44/100 (44%), Gaps = 14/100 (14%)
Query: 131 NEDTTSNTDETSNQNATSLDNS-TGMTANRNAY---VSLPQSEVNI---DVDNTTLRFAD 183
N + T TD T N N S DNS T + + N+ +S S+ N+ D +NT D
Sbjct: 313 NNNITITTDNT-NTNVISTDNSKTNVISKDNSNTHTISTDNSKTNVISTDNNNTDTISTD 371
Query: 184 NNTIDNGKTVNKSS-----NESNQNAKRNQNQKGNAKGT 217
N+ D T N ++ N +N + K N + K T
Sbjct: 372 NDNTDTKATDNDNTDTKATDNNNNTDTKATDNNNTDTKAT 411
 Score = 34.4 bits (77), Expect = 0.81
 Identities = 28/101 (27%), Positives = 41/101 (39%), Gaps = 15/101 (14%)
Query: 131 NEDITSNIDETSNQNATSLDNSTGMTA--NRNAYVSLPQSEVNIDV-----DNTTLRFA 182
N DT + ++ ++ AT +N+T A N N N D +NT +
Sbict: 374 NTDTKATDNDNTDTKATDNNNTDTKATDNNNTDTKATDKSNNTDTKATDNNNTDTKAT 433
Query: 183 DNNTIDNGK-----TVNKSSNESNQNAKRNQNQKGNAKGT 217
           DNN NK T K+++ +N NK N NK T
Sbjct: 434 DNNN-TNTKATDSNNTNTKATDNNNTNTKATDNNNTNTKAT 473
Score = 32.5 bits (72), Expect = 3.1
Identities = 30/110 (27%), Positives = 40/110 (36%), Gaps = 23/110 (20%)
Query: 131 NEDTTSNTDETSNQNATSLDNS-----TGMTANRNAYVSLPQS----EVNIDVDNTTLRF 181
N +TT N ++N S DN+ T T N N + + D NT ++
Sbjct: 251 NINTTQNLTTSTNTTTVSTDNNNNNINTKPTDNNNTNIKSTDNYNTGTKETDNKNTDIKA 310
Query: 182 ADNNTI-------DNGKTVNKSSNESNQNAKRNQNQKGNAKGT 217
DNN I DN KT S + SN + N K N T
Sbjct: 311 TDNNNITITTDNTNTNVISTDNSKTNVISKDNSNTHTISTDNSKTNVIST 360
>gi|1429240|emb|CAA67659| (X99260) lower collar protein
           [Bacteriophage B103]
           Length = 293
Score = 43.8 bits (101), Expect = 0.001
Identities = 53/204 (25%), Positives = 79/204 (37%), Gaps = 42/204 (20%)
Query: 56 EKVFKG----FSLKDELSDLLFKKSFTIHFLD----REINRQTVEAFGMQVITVCITHED 107
           EK+ KG F + + D ++K F HF+ REI +T F + T I +
Sbjct: 26 EKIEKGRPKLFDFQYPIFDESYRKVFETHFIRNFYMREIGFETEGLFKFNLETWLIINMP 85
Query: 108 YLNVVYSSSEVEKY------LQSQGFTEH-----NEDTT------SNTDETSN
Y N ++ S E+ KY L + G ++ N DTT SNT +
                                                             ---SNTDETSNQNA 146
Sbjct: 86 YFNKLFES-ELIKYDPLENTRLNTTGNKKNDTERNDNRDTTGSMKADGKSNTKTSDKTNA 144
Query: 147 TSLDNSTGMTA------NRNAYVSLPQSEVNIDVDN--TTLRFADNNTIDNGKTVNKS 196
T GT NR PS +N+ ++ TL +A + I + T NK
Sbjct: 145 TGSSKEDGKTTGSVTDDNFNRKIDSDQPDSRLNLTTNDGQGTLEYA--SAIEENNTNNKR 202
Query: 197 SNESNQNAKRNQNQKGNAKGTQFT 220
                                                                                   N + + GT T
Sbjct: 203 NTTGTNNVTSSAESESTGSGTSDT 226
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Query= pt|110879 44AHJDORF009 Phage 44AHJD ORF |5744-6496|2 1 (250 letters)

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>gi|2764981|emb|CAA69021.1| (Y07739) N-acetylmuramoyl-L-alanine
            amidase (Staphylococcus phage Twort)
            Length = 467
 Score = 180 bits (452), Expect = 1e-44
 Identities = 89/157 (56%), Positives = 109/157 (68%), Gaps = 8/157 (5%)
Query: 1 MKSQQQAKEWIYKHEGAGVDFDGAYGFQCMDLSVAYVYYITDGKVRMWGNAKDAINNDFK 60
            MK+ +QA+ +I G DFDG YG+QCMDL+V Y+Y++TDGK+RMWGNAKDAINN F
Sbjct: 1 MKTLKQAESYIKSKVNTGTDFDGLYGYQCMDLAVDYIYHVTDGKIRMWGNAKDAINNSFG 60
Query: 61 GLATVYKNTPSFKPQLGDVAVYTNGQ---YGHIQCVLS----GNLDYYTCLEQNWLGGGF 113 G ATVYKN P+F+P+ GDV V+T G YGHI V + G+L Y T LEQNW G G
Sbjct: 61 GTATVYKNYPAFRPKYGDVVVWTTGNFATYGHIAIVTNPDPYGDLQYVTVLEQNWNGNGI 120
Query: 114 DGWEKATIRTHYYDGVTHFIRPKFSGSNS-KALETSK 149
E ATIRTH Y G+THFIRP F+ +S K +T K
Sbjct: 121 YKTELATIRTHDYTGITHFIRPNFATESSVKKKDTKK 157
 Score = 61.7 bits (147), Expect = 6e-09
Identities = 41/125 (32%), Positives = 57/125 (44%), Gaps = 8/125 (6%)
Query: 125 YYDGVTHFIRPKFSGSNSKALETSKVNTFGKWKRNQYGTYYRNENGTFTC-GFLPIFARV 183
YY+G T P +K + +T G W N YGTYY++E+ TF C I R
Sbjct: 346 YYEGKTPV--PTVVNQKAKTKPVKQSSTSG-WNVNNYGTYYKSESATFKCTARQGIVTRY 402
Query: 184 GSPKLSEPNGYWFQPNGYTPYNEVCLSDGYVWIGYNWQGTR-YYLPVRQWNGKTGNSYSV 242
                                Y+ VC DGYVWI + G + ++PVR W+ N+
Sbjct: 403 TGPFTTCPQAGVLYYGQSVTYDTVCKQDGYVWISWTTNGGQDVWMPVRTWD---KNTDIM 459
Query: 243 GIPWG 247
Sbjct: 460 GQLWG 464
>gi|113675|sp|P24556|ALYS_STAAU AUTOLYSIN
            (N-ACETYLMURAMOYL-L-ALANINE AMIDASE)
            >gi|79887|pir||JQ1147 N-acetylmuramoyl-L-alanine amidase
            (EC 3.5.1.28) - Staphylococcus aureus >gi 153067
            (M76714) peptidoglycan hydrolase [Staphylococcus aureus]
            Length = 481
 Score = 118 bits (292), Expect = 6e-26
 Identities = 56/117 (47%), Positives = 68/117 (57%), Gaps = 1/117 (0%)
Query: 135 PKFSGSNSKALETSKVNTFGK-WKRNQYGTYYRNENGTFTCGFLPIFARVGSPKLSEPNG 193
            P + SN + ++ V WKRN+YGTYY E+ FT G PI R P LS P G
Sbjct: 365 PVATVSNESSASSNTVKPVASAWKRNKYGTYYMEESARFTNGNQPITVRKVGPFLSCPVG 424
Query: 194 YWFQPNGYTPYNEVCLSDGYVWIGYNWQGTRYYLPVRQWNGKTGNSYSVGIPWGVFS 250
            Y FQP GY Y EV L DG+VW+GY W+G RYYLP+R WNG + +G WG S
Sbjct: 425 YQFQPGGYCDYTEVMLQDGHVWVGYTWEGQRYYLPIRTWNGSAPPNQILGDLWGEIS 481
 Score = 78.0 bits (189), Expect = 7e-14
 Identities = 48/109 (44%), Positives = 62/109 (56%), Gaps = 6/109 (5%)
Query: 15 EGAGVDFDGAYGFQCMDLSVAYVYYITDGKVRMWGNAKDA-INNDFKGLATVYKNTPSFK 73
                + D YGFOC D + A + + G + AKD N+F GLATVY+NTP F
Sbjct: 18 EGKQFNVDLWYGFQCFDYANAG-WKVLFGLLLKGLGAKDIPFANNFDGLATVYQNTPDFL 76
Query: 74 PQLGDVAVYTNGQ---YGHIQCVLSGNLDYYTCLEQNWLGGGF-DGWEK 118
                          YGH+ V+ LDY EQNWLGGG+ DG E+
Sbjct: 77 AQPGDMVVFGSNYGAGYGHVAWVIEATLDYIIVYEQNWLGGGWTDGIEQ 125
>gi|1763243 (U72397) amidase [bacteriophage 80 alpha]
           Length = 481
                                                                                    Score = 118 bits (292), Expect = 6e-26
 Identities = 56/117 (47%), Positives = 68/117 (57%), Gaps = 1/117 (0%)
Query: 135 PKFSGSNSKALETSKVNTFGK-WKRNQYGTYYRNENGTFTCGFLPIFARVGSPKLSEPNG 193
P + SN + ++ V WKRN+YGTYY E+ FT G PI R P LS P G
Sbjct: 365 PVATVSNESSASSNTVKPVASAWKRNKYGTYYMEESARFTNGNQPITVRKVGPFLSCPVG 424
```

Query: 194 YWPQPNGYTPYNEVCLSDGYVWIGYNWQGTRYYLPVRQWNGKTGNSYSVGIPWGVFS 250

```
Y FQP GY Y EV L DG+VW+GY W+G RYYLP+R WNG + +G WG S
Sbjct: 425 YQPQPGGYCDYTEVMLQDGHVWVGYTWEGQRYYLPIRTWNGSAPPNQILGDLWGEIS 481
 Score = 83.5 bits (203), Expect = 2e-15
 Identities = 50/115 (43%), Positives = 65/115 (56%), Gaps = 6/115 (5%)
          EWIYKHEGAGVDFDGAYGFQCMDLSVAYVYYITDGKVRMWGNAKDA-INNDFKGLATVYK 67
           EW+ EG + D YGFQC D + A + + G + AKD N+F GLATVY+
Sbjct: 12 EWLKTSEGKQFNVDLWYGFQCFDYANAG-WKVLFGLLLKGLGAKDIPFANNFDGLATVYO 70
Query: 68 NTPSFKPQLGDVAVYTNGQ---YGHIQCVLSGNLDYYTCLEQNWLGGGF-DGWEK 118
NTP F Q GD+ V+ + YGH+ V+ LDY EQNWLGGG+ DG E+
           NTP F Q GD+ V+ +
Sbjct: 71 NTPDFLAQPGDMVVFGSNYGAGYGHVAWVIEATLDYIIVYEQNWLGGGWTDGIEQ 125
>gi|4574237|gb|AAD23962.1|AF106851_1 (AF106851) LytN [Staphylococcus
           aureusl
           Length = 383
 Score = 84.3 bits (205), Expect = 9e-16
 Identities = 48/128 (37%), Positives = 68/128 (52%), Gaps = 7/128 (5%)
Query: 15 EGAGVDFDGAYGFQCMDLSVAYVYYITDGKVRMWGNAKDAINNDFKGLATVYKNTPSFKP 74
           E G DFDG+YG+QC DL Y ++ +++G
                                                      N+F A +Y NTP+FK
Sbjct: 252 ENRGWDFDGSYGWQCFDLVNVYWNHLYGHGLKGYGAKDIPYANNFNSEAKIYHNTPTFKA 311
Query: 75 QLGDVAVYT---NGQYGHIQCVLSGNLD----YYTCLEQNWLGGGFDGWEKATIRTHYYD 127
+ GD+ V++ G YGH VL+G+ D + L+QNW GG+ E A H Y+
Sbjct: 312 EPGDLVVFSGRFGGGYGHTAIVLNGDYDGKLMKFQSLDQNWNNGGWRKAEVAHKVVHNYE 371
Query: 128 GVTHFIRP 135
               FTRP
Sbjct: 372 NDMIFIRP 379
gi|3767593|dbj|BAA33856.1| (AB015195) LytN [Staphylococcus aureus]
           Length = 383
Score = 84.3 bits (205), Expect = 9e-16
Identities = 48/128 (37%), Positives = 68/128 (52%), Gaps = 7/128 (5%)
Query: 15 EGAGVDFDGAYGFQCMDLSVAYVYYITDGKVRMWGNAKDAINNDFKGLATVYKNTPSFKP 74
           E G DFDG+YG+QC DL Y ++ +++G
                                                     N+F A +Y NTP+FK
Sbjct: 252 ENRGWDFDGSYGWQCFDLVNVYWNHLYGHGLKGYGAKDIPYANNFNSEAKIYHNTPTFKA 311
Query: 75 QLGDVAVYT---NGQYGHIQCVLSGNLD----YYTCLEQNWLGGGFDGWEKATIRTHYYD 127 + GD+ V++ G YGH VL+G+ D + L+QNW GG+ E A H Y+
Sbjct: 312 EPGDLVVFSGRFGGGYGHTAIVLNGDYDGKLMKFQSLDQNWNNGGWRKAEVAHKVVHNYE 371
Query: 128 GVTHFIRP 135
               FIRD
Sbjct: 372 NDMIFIRP 379
>gi|2764983|emb|CAA69022.1| (Y07740) cell wall hydrolase Ply187
           [Staphylococcus phage 187]
           Length = 628
 Score = 76.9 bits (186), Expect = 2e-13
 Identities = 50/144 (34%), Positives = 68/144 (46%), Gaps = 18/144 (12%)
Query: 5 QQAKEWIYKHEGAGVDFDGAYGFQCMDLSVAYVYYITDGKVRMW-----GNAKDAINNDF 59
           +Q +W G+GVD DG YG QC DL Y++ R W GNA+D
Sbjct: 12 KQVVDWAINLIGSGVDVDGYYGRQCWDLP-NYIFN-----RYWNFKTPGNARDMAWYRY 64
Query: 60 KGLATVYKNTPSFKPQLGDVAVYTNGQY----GHIQCVLS-GNLDYYTCLEQNWLGGGF 113
               V++NT F P+ GD+AV+T G Y GH V+ Y+ ++QNW
Sbjct: 65 PEGFKVFRNTSDFVPKPGDIAVWTGGNYNWNTWGHTGIVVGPSTKSYFYSVDQNWNNSNS 124
Query: 114 DGWEKATIRTHYYDGVTHFIRPKF 137
               A H Y GVTHF+RP
Sbict: 125 YVGSPAAKIKHSYFGVTHFVRPAY 148
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>gi|3287732|sp|005156|ALE1_STACP GLYCYL-GLYCINE ENDOPEPTIDASE ALE-1
             PRECURSOR >gi|1890068|dbj|BAA13069| (D86328) ALE-1
             (Staphylococcus capitis)
             Length = 362
 Score = 73.4 bits (177), Expect = 2e-12
 Identities = 47/117 (40%), Positives = 61/117 (51%), Gaps = 10/117 (8%)
Query: 132 FIRPKFSGSNSKALETSKVNTFGKWKRNQYGTYYRNENGTFTCGFLPIFARVGSPKLSEP 191
F++ GSNS TS N G +K N+YGT Y++E+ +FT I R+ P S P
Sbjct: 252 FLKSAGYGSNS----TSSSNNNG-YKTNKYGTLYKSESASFTAN-TDIITRLTGPFRSMP 305
Query: 192 NGYWFQPNGYTPYNEVCLSDGYVWIGYNW-QGTRYYLPVRQWNGKTGNSYSVGIPWG 247
+ Y+EV DG+VW+GYN G R YLPVR WN TG +G WG
Sbjct: 306 QSGVLRKGLTIKYDEVMKQDGHVWVGYNTNSGKRVYLPVRTWNESTG---ELGPLWG 359
>gi|79926|pir||A25881 lysostaphin precursor - Staphylococcus
             simulans >gi | 153047 (M15686) lysostaphin (ttg start
             codon) [Staphylococcus simulans]
             Length = 389
 Score = 69.5 bits (167), Expect = 3e-11
 Identities = 48/133 (36%), Positives = 62/133 (46%), Gaps = 20/133 (15%)
Query: 131 HFIRPKFSGSNSKALETS---KVNTFGK------------WKRNQYGTYYRNENGTFTCG 175
HF R S SNS A + K +GK WK N+YGT Y++E+ +FT
Sbjct: 258 HFQRMVNSFSNSTAQDPMPFLKSAGYGKAGGTVTPTPNTGWKTNKYGTLYKSESASFTPN 317
Query: 176 FLPIFARVGSPKLSEPNGYWFQPNGYTPYNEVCLSDGYVWIGYNW-QGTRYYLPVRQWNG 234
I R P S P + Y+EV DG+VW+GY G R YLPVR WN
Sbjct: 318 -TDIITRTTGPFRSMPQSGVLKAGQTIHYDEVMKQDGHVWVGYTGNSGQRIYLPVRTWNK 376
Query: 235 KTGNSYSVGIPWG 247
T ++G+ WG
Sbjct: 377 STN---TLGVLWG 386
>gi|126496|sp|P10548|LSTP_STAST LYSOSTAPHIN PRECURSOR
             (GLYCYL-GLYCINE ENDOPEPTIDASE) >gi|79927|pir||S01079
            lysostaphin precursor - Staphylococcus simulans bv.
            staphylolyticus >gi|581744|emb|CAA29494| (X06121)
            lysostaphin (AA 1-480) [Staphylococcus simulans bv.
            staphylolyticus]
            Length = 480
 Score = 69.5 bits (167), Expect = 3e-11
 Identities = 48/133 (36%), Positives = 62/133 (46%), Gaps = 20/133 (15%)
Query: 131 HFIRPKFSGSNSKALETS---KVNTFGK------------WKRNQYGTYYRNENGTFTCG 175
HF R S SNS A + K +GK WK N+YGT Y++E+ +FT
Sbjct: 349 HFQRMVNSFSNSTAQDPMPFLKSAGYGKAGGTVTPTPNTGWKTNKYGTLYKSESASPTPN 408
Query: 176 FLPIFARVGSPKLSEPNGYWFQPNGYTPYNEVCLSDGYVWIGYNW-QGTRYYLPVRQWNG 234
I R P S P + Y+EV DG+VW+GY G R YLPVR WN
Sbjct: 409 -TDIITRTTGPFRSMPQSGVLKAGQTIHYDEVMKQDGHVWVGYTGNSGQRIYLPVRTWNK 467
Query: 235 KTGNSYSVGIPWG 247
T ++G+ WG
Sbjct: 468 STN---TLGVLWG 477
lysostaphin (Staphylococcus simulans)
            Length = 493
 Score = 69.5 bits (167), Expect = 3e-11
Identities = 48/133 (36%), Positives = 62/133 (46%), Gaps = 20/133 (15%)
Query: 131 HFIRPKFSGSNSKALETS---KVNTFGK------WKRNQYGTYYRNENGTFTCG 175
                                K +GK
            HFR S SNS A +
                                                         WK N+YGT Y++E+ +FT
Sbjct: 362 HFQRMVNSFSNSTAQDPMPFLKSAGYGKAGGTVTPTPNTGWKTNKYGTLYKSESASFTPN 421
Query: 176 FLP1FARVGSPKLSEPNGYWFQPNGYTPYNEVCLSDGYVW1GYNW-QGTRYYLPVRQWNG 234
```

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I R P S P
                                           Y+EV DG+VW+GY G R YLPVR WN
 Sbjct: 422 -TDIITRTTGPFRSMPQSGVLKAGQTIHYDEVMKQDGHVWVGYTGNSGQRIYLPVRTWNK 480
Query: 235 KTGNSYSVGIPWG 247
                ++G+ WG
Sbjct: 481 STN---TLGVLWG 490
 >gi|3341932|dbj|BAA31898.1| (AB009866) amidase (peptidoglycan
            hydrolase) [bacteriophage phi PVL]
             Length = 484
 Score = 68.3 bits (164), Expect = 6e-11
 Identities = 52/150 (34%), Positives = 71/150 (46%), Gaps = 17/150 (11%)
            SQQQAKEWIYKHEGAGVDFDGAYGFQCMDLSVAYVYYITDGKVRMWGNAKDAINNDFKGL 62
            ++ QA++W G + D YGFQC D + + + I G+ R+ G I D K
TKNQAEKWFDNSLGKQFNPDLFYGFQCYDYASMF-FMIATGE-RLQGLYAYNIPFDNKAR 61
Sbict: 4
Query: 63 ATVY----KNTPSFKPQLGDVAVYTN---GQYGHIQCVLSGNLDYYTCLEQNWLGGGF-- 113
Y KN SF PQ D+ V+ + G GH++ V S NL+ +T QNW G G+
Sbjct: 62 IEKYGQIIKNYDSFLPQKLDIVVFPSKYGGGAGHVEIVESANLNTFTSFGQNWNGKGWTN 121
Query: 114 ----DGW--EKATIRTHYYDGVTHFIRPKF 137
GW E T HYYD +FIR F
Sbjct: 122 GVAQPGWGPETVTRHVHYYDDPMYFIRLNF 151
Query= pt | 110882 44AHJDORF012 Phage 44AHJD ORF | 8391-8813 | 3 1
           (140 letters)
>gi|140528|sp|P24811|YQXH_BACSU HYPOTHETICAL 15.7 KD PROTEIN IN
            SPOILIC-CWLA INTERGENIC REGION (ORF2)
            >gi|322189|pir||B44816 orf2 5'of autolytic amidase -
            Bacillus subtilis >gi|142801 (M59232) open reading frame
2 [Bacillus subtilis] >gi|1217874|dbj|BAA06959| (D32216)
ORF121 [Bacillus subtilis] >gi|1303767|dbj|BAA12423|
             (D84432) YqdD [Bacillus subtilis]
            >gi|2635036|emb|CAB14532| (Z99117) alternate gene name:
            yqdD; similar to holin (Bacillus subtilis)
            Length = 140
 Score = 80.4 bits (195), Expect = 6e-15
 Identities = 45/130 (34%), Positives = 67/130 (50%), Gaps = 3/130 (2%)
           VKFRFTDSEAFHMFIYAGDLKLLYFLFVLMFVDIITGISKAIKNNNLWSKKSMRGFSKKX 63
            + F D ++F G +K L LVL +D++TG+ KA K L S+ + G+ +K
INFETLDLARVYLF---GGVKYLDLLLVLSIIDVLTGVIKAWKFKKLRSRSAWFGYVRKL 64
Query: 64 XXXXXXXXXXXXXXXXXXKGGLLMITIFYYIANEGLSIVENCAEMDVLVPEQIKDKLRVI 123
                                 G L T+ +YIANEGLSI EN A++ V +P I D+L+ I
Sbjct: 65 LNFFAVILANVIDTVLNLNGVLTFGTVLFYIANEGLSITENLAQIGVKIPSSITDRLQTI 124
Query: 124 KNDTEKSDNN 133
+N+ E+S NN
Sbjct: 125 ENEKEQSKNN 134
>gi|4126631|dbj|BAA36651.1| (AB016282) ORF45 (bacteriophage phi-105)
            Length = 135
 Score = 76.1 bits (184), Expect = 1e-13
 Identities = 44/115 (38%), Positives = 61/115 (52%), Gaps = 4/115 (3%)
G++K L + VL +DIITG+ KA K L S+ + G+ +K
Sbjct: 17 GEVKYLDLMLVLNIIDIITGVIKAWKFKELRSRSAWFGYVRKMLSFLVVIVANAIDTIMD 76
Query: 81 XKGGLLMITIFYYIANEGLSIVENCAEMDVLVPEQIKDKLRVIKND----TEKSD 131
                                                                                           G L T+ +YIANEGLSI EN A++ V +P I D+L VI++D TEK D
Sbjct: 77 LNGVLTFATVLFYIANEGLSITENLAQIGVKIPAVITDRLHVIESDNDQKTEKDD 131
>gi|141088|sp|P26835|YNGD_CLOPE HYPOTHETICAL 14.9 KD PROTEIN IN NAGH
            3'REGION (ORFD) >gi|1075967|pir||S43905 hypothetical
            protein D - Clostridium perfringens >gi|455154 (M81878)
```

311

ORF D [Clostridium perfringens] Length = 132

Score = 60.9 bits (145), Expect = 4e-09 Identities = 38/127 (29%), Positives = 63/127 (48%), Gaps = 3/127 (2%)

MNEVKFRFTDSEAFHMFIY-AGDLKLLYFLFVLMFVDIITGISKAIKNNNLWSKKSMRGF 59

+I+ A D+ L+ L V +F+D +TG+ K K+ L S INYIKWGIVSLGTLFTWIFGAWDIPLITLL-VFIFLDYLTGVIKGCKSKELCSNIGLRGI 63 Sbict: 5

Query: 60 SKKXXXXXXXXXXXXXXXXXXXXXXXXXIII-FYYIANEGLSIVENCAEMDVLVPEQIKD 118

+KK + I ++YI NEG+SI+ENCA + V +PE++K
Sbjct: 64 TKKGLILVVLLVAVMLDRLLDNGTWMFRTLIAYFYIMNEGISILENCAALGVPIPEKLKQ 123

Query: 119 KLRVIKN 125 Sbjct: 124 ALKQLNN 130

amidase (Bacillus subtilis) Length = 134

Score = 36.4 bits (82), Expect = 0.099 Identities = 25/109 (22%), Positives = 41/109 (36%)

Query: 17 FIYAGDLKLLYFLFVLMFVDIITGISKAIKNNNLWSKKSMRGFSKKXXXXXXXXXXXXX 76

F + G L LM ++ I+ K + L KK KK
Sbjct: 20 FFFGGFQYSFLILLSLMAIEFISTTLKETIIHKLSFKKVFARLVKKLVTLALISVCHFFD 79

Query: 77 XXXXXKGGLLMITIFYYIANEGLSIVENCAEMDVLVPEQIKDKLRVIKN 125 +G + + I +YI E + IV + + + VP+ + D L +KN

Sbjct: 80 QLLNTQGSIRDLAIMFYILYESVQIVVTASSLGIPVPQMLVDLLETLKN 128

>gi|1181973|emb|CAA87743.1| (Z47794) holin protein (Bacteriophage CP-1 Length = 134

Score = 31.3 bits (69), Expect = 3.3 Identities = 27/88 (30%), Positives = 36/88 (40%), Gaps = 5/88 (5%)

LF L+ D ITG KA K S ++G K G +L
Sbjct: 18 LFALILFDFITGFLKAWKWKVTDSWTGLKGVIKHTLTFIFYYFVAVFLTYIHAMAVGQIL 77

Query: 87 MITIFYYIANEGLSIVENCAEMDVLVPE 114

++ I Y A LSI+EN A M V +P+
Sbjct: 78 LVIINLYYA---LSIMENLAVMGVFIPK 102

Table 21

# Phage 182 complete genome sequence. 17833 nucleotides.

1	tagaatattg	tcataaaaca	caaacataat	aatgcatatt	attgtttaca	aatatgtaat	ttcgtgatat
71						ttagaaattg	
141							cagaattaga
211 281	atotttato	atgaactaga	ggaattggtg	gacgaagtaa	acgatattgc	taaagatccg	gaggaaagat
351	ttacaacgaa	aagatcacaa	atctgaaatc	ggaaatcata	tractorasa	tetgeaagat	aaaactagta
421	aaataattac	acaaaaagct	ttacaaatat	aacacatcat	ottatactaa	aagagtagta	addactagta
491						gaaaataatg	
561	taatgcaacg	aaacgtaaca	tcaactaaag	tagaattctc	agaagttatc	gtacaagatg	gagegeeaac
631	aattgtacca	tgcgaaccag	ttgtcttaac	aggaaaactt	tcagaagaaa	aagctttatc	agcgatcaaa
701	cgtaaaaacc	ctgataaaaa	cgtagttgta	acaaatgttt	cacatgaaac	agcgctttac	acaatgccag
771						caaaactaaa	
841 911						cagaaggaat	
981						agtactagaa	
1051	cttcacagaa	gatggtaaaa	cttatocooo	totatoaca	cyaaccacca	caagcagaac aatcagctaa	tagtcgcttt
1121	gatatgatga	ctoctaaccc	tgacatcaaa	ccaaaaattt	cttttgtcga	aggaaaatca	aaacctaatt
1191	aaaaatttgt	aaatctacaa	gtggtttcac	tgtagcataa	aaatacagga	atctagtaag	ccacttagcg
1261						tgaccgtaag	
1331	aatgatagag	ccaagttaga	gaaaatctac	ggtaaatcta	acaaagctcg	taaaaaatac	aatcgtttaa
1401	gacaaaaagg	agttgaggaa	aggcaacttc	caactgttcc	aacatcaaag	aaaagactta	ttgactacgt
1471	aaaatcaaca	aatatgagtc	gtagtgattt	taacaagatg	ttagacgagt	tggtagattt	tgcacaacct
1541 1611	tacaacgaga	attacatttt	tgagatcaac	aagcgaaatg	ttgcaatctc	aagagcgcaa	atcaaagaag
1681	taagaaggg	aacagagcaa	geccaaaaag	cgaaagaaga	acactacaaa	gagettaaca taggtgetga	aagttgaagt
1751						tcagtcttat	
1821	taggaaaaca	agacgaacaa	tattttgacg	aaagagacca	actttattac	gacaatttca	cagaaaata
1891	gtttactatt	ttcaattcag	acgctgacga	tattgttcgt	ttacttgact	caatggggct	tgatctattt
1961	atgaaaacat	atgttagtaa	cttcttagac	atgaaccttg	actacattta	tgacgaagca	gaagtacaac
2031	agaaaaaaga	acaagtttac	agtaagattg	caaaagtgat	cgagtctgaa	acaggtggag	aagtcccctc
2101	atataacccc	acgaagaaca	tcacaattaa	ttcagaaaca	ggagaagaat	tatgattaag	aaatatactg
2171 2241	gcgactttga	aacaacaact	gatctcaacg	attgtcgtgt	atggtcgtgg	ggcgtatgcg	atatagacaa
2311	cgttgacaat	atgacgttcg	gtttagaaat	cgattcttt	tttgagtggt	gtaaaatgca	aggcagcaca
2381	aatgototaa	agaaggaaaa	gaagatcgaa	catteteese	acticates	gttattcaaa aatatgggtc	aatggtttca
2451						cgaaaaaaga	
2521						tgcagaagct	
2591	ctataaaaaa	aggcgaaata	gattatacaa	aagaaagacc	tattggttac	aaaccaacaa	aagatgaatg
2661	ggagtattta	aagaacgaca	ttcagattat	ggcgatggca	ttaaaaattc	aattcgatca	aggactaact
2731	cgaatgacta	gaggaagcga	cgctttaggc	gattacaaag	attggctaaa	agctacacat	ggaaaatcaa
2801						cgtaaagcat	
2871 2941						ttgtctttga	
3011	aaccgaacaa	ccaaatgta	ctgtagacct	assistances	gaacacctct	attctacgaa cgtttaaagg	ggagaataca
3081	tccaaccatt	caagttaagc	aaagttcatt	attcattcaa	aacgaatatc	ttgaatcaag	totaaacaac
3151	ttaggagttg	acqaattaat	cgatcttact	cttacaaatq	ttgacctaga	attattttt	gaacactacg
3221	atattttaga	gatacattac	acttacggat	atatgttcaa	agcttcttgt	gatatgttca	aaggctggat
3291	cgataaatgg	atcgaagtaa	agaacaccac	cgaaggggct	agaaaagcta	acgccaaagg	tatgttaaat
3361	agcttgtatg	gaaagttcgg	aacaaaccct	gacattacag	gaaaagtgcc	ttacatgggc	gaggacggca
3431 3501	ttgttcgatt	gacactagga	gaagaagaat	taagagatcc	tgtttatgtt	ccgcttgcta	gttttgtgac
3571						gcattattta	
3641	gttattocoo	gcatgaaaor	acatttcaac	gaggaaaatt	cattengean	ggttgatcct aaaacatacg	tagaagaaat
3711	tgatggcgaa	ttaaatgtaa	agtgtgctgg	tatoccapat	caataaaaa	agattgtaac	ttttaacaat
3781	tttgaagttg	gtttttcaag	ctatggaaag	ttqctaccta	aaagaacaca	aggtggcgtg	gtattagtag
3851	acacaatgtt	tacaatcaaa	taaggaggac	taataatgga	actatataaa	gcaatgttta	togtacgtga
3921	tgaaggtact	attgacggtt	acgatactga	acactatgta	gatatttctt	tacatgactt	tgaagaaata
3991	tatggaaaag	aaacacgtga	aattgaagca	gtaacattag	taaaaacaqq	aaatttaaaa	aaataaatta
4061	tttacatcct	ttgcaaagta	tggtaaaata	ttcttgtgat	agttgacaag	agtcaaattt	ggcgagattg
4131 4201	ggcgaatgta	cacgtgaaat	accgtgcgct	cccgttaagt	tatggacaca	taaacgtttt	gaccgtcaac
4201	ttcacoross	acacttttag	tootateete	caaacgtggc	cactetttt	tgtgtttcac	agaattatgt
4341	qaatcaattt	taaatootat	tcttgaaagr	gtcacacaca	gagguggaga	ttatggaaat atcaaagatt	caaagaacat
4411	ttgaaqcatt	gcgagaagac	tacqqaqcaa	caactgaage	tttgacatca	gcaaatagca	yeayaacatt cacttoaaaa
4481	gttaaagaaa	gataacgaaq	cgttggttat	ttcaaactca	aaattgttcc	gagaacgagc	gatcgtagaa
4551	ccagcagaaa	ataacgaacc	agaaacagac	cagaatatta	cactagacga	tttaqqaatt	taaqqaqqaa
4621	aaaacatggc	tgacaaaatc	acagaacaag	atgttcttcg	tgccacaaat	gtagaaacac	cagtacaatt
4691	aatgactgct	atttataata	gttcatcatc	tctttttcag	gcgaacgtac	ctatgccaaa	tgcagataac

atcgaagcgg ttggtgcagg gatcacacgt ttagacgtag taaaaaacga atttatttca actttagttg accgtattgg taaagtagtt atccgataca aatcttggcg taaccctttg aaaatgttta aaaaaggaaa 4761 4831 4901 catgccttta ggtcgaacga ttgaagaaat ttttgttgac attgcacaqq aacataaqtt caaccctqac gagtotgtta caggggtatt taaacaggaa gttcccgatg taaaaacatt gttccacgaa attaatcgtg 4971 aaggttacta caaacaaacg atccaagaag catggttaga aaaagcattt acttcatggg ataatttcaa 5041 tagtttcgtt gctggtgtaa tgaacgcttt atacacaggt gacgaagtaa gcgaatttga atacacgaaa 5111 5181 ttattaatag caaactacca agaaaaagag ctattcaaag agatcgaaat tqqcqaaatt actqaatcaa 5251 atgcaaaaga atttatccgt aagatcaaat caacctctaa caaattagaa tttatgagtt ccgcttacaa cgctcaagga gttaaaacat ctacctcaaa atctgatcaa tacgttatta ttgacgccga cacagacgca 5321 5391 accattgacg ttgacgtttt agcagcggca ttcaatatga gtaaaactga ctttgtagga cacaaaatcg ttattgatga gtttcctaaa aaagaaggcg aagaatcgtc aaatattgtg gcagttattg tagatagtga 5461 atggtttatg atctacgaca aattgtacaa aacaacaagt ctatacaacc ctgaagggtt atattggaat 5531 tattggttgc accaccacca actatattct acttctcaat tcgggaacgc tgttgctttt gttaaatcag 5601 caacaaaacc tgtcacaaaa gttgcttttg caagtgcaac aactagtgtt gttaaaggat catctaaaga 5671 5741 tatcgcattg acatttacac cagtagaagc aacaaaccaa caaggagaag ttgtttcatc agcaccagca 5811 ttggttaagg caaccgtaaa acaaacagca ggtaaagcga ctgccgtaac cgtagaaggc ttagaagtcg 5881 gtcaatcatt agtaacattc acagctatcg gaggtcaaca agcaacggtt cttgttacgg ttacttctga 5951 ctaaggagga caattatggc aagaaggtat acaaatgtaa aattgtiggc taacgtgcct tttgataaca cctatacaca cacaagatgg tttaaaactc aacaggaaca ggaatcgtac tttaattcgt ttcctgttct 6021 taacgagaat agagattgtt cttatcaaag ggatacacaa ctcgggggag tttttagagt agataaacac 6091 aaagacgcct tatatgcttg taactatctc atctttaaaa acgaagaaac ttatcctagt aaatggcagt atgcctttgt tactgatatt gaatataaga atgacaacac aagtttcgtt acctttgaaa ttgatgtttt 6161 6231 acaaacttat cytttcgata ttggtatacg agaaagtttc attgcaaaag acaccctca actttattat tcgaatggaa tacctttcat taatacaatt gaagagtcgc ttgattacgg tagagaatac acaacaacaa 6301 6371 atgtaacaac ttttcatcct aacgatggag tcaattttct tgttattcta acaagtgaag caatgccagt 6441 tggagataag gaagataaat caggaggatc aatagtaggt ggcccatctc ctttttccta ttatttactt 6511 cctatcaatt caagtgggga ggtatacaaa ccaaatgggg caggcaatgc taattttgga gagtacatgg 6581 cgtttcttac aacgaaagaa ccttttttaa ataagatagt cgggatgtat gtaacgtcgt atacaggtat 6651 accattcatt gtggatcacg cgaacaaaac ggtaaggtat aatgcaggag gttcttataa gatcatqctt 6721 6791 ccaacctacg ctagtgatcc aacaggaaca atgaaaacat tcgctttctt ttgtgtaaaa gaagcaagaa 6861 cattegtace taaaagaatt gatettgtag ggaaegtgta taactaettt agagaagett tteegtttaa 6931 tgttaaggaa tcaaaactat ttatgtatcc ctattgttta atagaaatta cagatacaaa aggacatgta 7001 atgactttaa gacctgaata tottacaggt ggtaaattga gtgtatatgt aaaaggttog ttaggaattt 7071 ctaataaagt gatgatcgag ccgattgatt atgatgtaag taactcaacc attattacca atttaagtga 7141 caagatgtta atcgataatg atcctaacga tgtaggagtt aaatctgact atgcttctgc attcatgcaa 7211 ggaaacaaaa actccttgat tgctcaagag caaaacattc gcaatacttt cagacatggt atgggaaaca 7281 gtgcaatgag tacaggagga gcgatctttt cagccttagc aagtaacaac ccttttgttg gtttgactaa 7351 catcatggga gcaggacaac aagtaaacaa ctatgtttct gaaaaagaaa acggtttgaa cctcttggca ggtaaagtgg cagatatcga aaatattcca gataatgtaa cacagcttgg atcaaactta tctttcacaa 7421 caggaaactt tcaaaactat tatcaattgc gcttcaaaca aattaaatat gagtatgcaa caagacttga 7491 togttactto toaatgtatg goacaaagag caatogagta gotacaccaa acttacaaac aagaaaagca 7561 teggaattica ttaaattaaa agaaccaaat attgtaggca caatgagtaa cgatgtatta acacgtgtga aacaaattit tagtgcaggc gttacgcttt ggcatacgaa tgatgtttg aattataacc aagacaacgg 7631 7701 agatgtatag gaaggaggaa taagatgagt agacgaaaag gtgcaggact tgctagaaat aaccgttata cagcaaaaag cagaccttat ccaaatgaac cctattcaag tgatgtagaa gaaatcagct actatgaaca 7771 7841 7911 ttatcgtaga caactcacgc tccttacgtt tcagttgttt gaatgggaaa atttgccaaa atcaattgac 7981 cotogttatt tagaaattgc tttacacact aatggttatc ttggtttctt taaagaccct acacttgggt 8051 tcatggtttg cgcaggggca gaagatggtc aaatcgatca ttatcacaac cctattttct ttacagcaaa 8121 cgaagcaatg tatcacaaga gatatcctgt tttaagatat gatgatgatg atgataaatc aaaatgtatc 8191 atgttgtata ataatgactt gaaagtteet aegttaceaa gtttacateg ttttgettta gatatggegg 8261 acataaacca gatatcacga gtgaatcgaa gagcgcaaaa aacacctgta attattcaaa ctgatgaaaa 8331 gaaatacttc tcattgctac aagcttataa ccaaattgac gaaaataatc aggctgtttt tgtggataaa 8401 gatatggagt ttgacgaatc ttttaatgta tggcaaacaa atgctccata tgtagtagat aaactacgat 8471 Cagaattgaa cgaagtatgg aatgaagtgt taacttttct aggtatcaac aatgctaacg tagataagac 8541 tgcacgtgta caaacatcag aagtcttatc taacaatgaa cagattgaaa gttcaggtaa catcttgtta 8611 aaatcaagaa aagagttttg cgatcgtgta aatcgtgtct ttggcgatga acttgacgga aagattgacg tgaagtttag aacagacgcc gttcgacaat tacaactggc ggcaggtcaa tcaaaaaaag accagatgag tggagggttg ccaagtgcta cttaaacgtt atattgaaag tttcacttat taccaacctg aattatctcg 8681 8751 aaaagaacgt attgaagttg gccgaaaaca attgtttgat tttgattatc cgttttatga cgaacaaaaa cgagcagaat ttgaaacaaa atttatcaat cacttttact tgagagagat aggctcagaa acgatgggat 8821 8891 catttaagtt taatcttgac gaatatttaa atctaaacat gccctattgg aataaaatgt tcctatcaaa tcttgaagag tttccgattt ttgatgacat ggactacacc attgatgaga aacagaaatt gttaaatgag 8961 9031 attgatacaa acatcaaagc gaatcgtgat gaatcgaaga accaaacgaa gcaagtagat caaacagaca 9101 9171 acagaaacaa aaatacacgt gacacaggaa caaccgattc tttctcaagg aacacttata cagacacccc 9241 tcaaaaagat ttgagaattg ccagcaatgg agatggaaca ggtgtaatca attatgcaac aaatatcaca 9311 gaagatttga gtaaagaaac aacaagctcc acaggcgttg aaacaaacaa cgacaaaaca aatcaaaata 9381 Cacgaagcaa tgcttctgaa aaagaaacaa agaacacaga cattaataaa gatcaaaatc aaaccaaaga 9451 tacgattaca cgatataaag gtaaaaaggg aaacactgat tatgctgact tactcgaaaa atatcgtaga 9521 agtgttttga gaattgagaa aatgatcttt agagaaatga acaaggaagg cttatttctc cttgtttatg gagggaggta gcaacaatgg tagattttaa ccccgacaag cggtttgacg gtttacccgc tgtattcaaa --9591 9661 gaacgettta geaaatatee teatactgaa tacagatatg aattactatt agatgaagaa gtateggett 9731 taattgccta totgaatgaa gttggtgctt tagttaatga tatgagtggt tatttaaatt actttatcga 9801 acattttgtt gagaagttag aagagatcac aaatgacaca ctcaaaaaat ggttgtctga tggtacgtta 9871 gaaaatttaa tcaatgatac tgtttttgca aattatatca aagaaatcaa aagattacaa atcttggttg 9941 ctgaaacacg tgctaacagt gtgaatatte ttttgacaaa aaataaaccg gatgttgctg atgatcgaac

attitiggtat aaqaticaac gcgacaatac tgattatgga gccgatccta tigacacqtt acqtatiqti

314

10081 gcaatcaata aagttagtgg ctggaatacc gctacaggag atatttatct taacattaaa ggaacggagg gtgtataatg gcagacatta gaacacact aacaagtgaa gatggatcag acaatttatt tccaatttca aaagccgtta atattatgac taatagcggt acgaatgtag aaggagaatt gggtacactc aaacaaaatg 10151 10221 10291 acgaaacaat gaatacctca gttcaaaatg ctgtagttac tgccaatcaa gcaaaagatt ctgtagctga attaaatgta aatgttggta aactaaccaa tcgaataaca acattagaga gtacagtggc taatcttgat 10361 ggtattegtt atgtagaggt gtaatatgge agataahaat atteaahtge aggatahaaga teataategt ttaatgeetg ttacaattgt ttaatgeetg ttaatgetgaaa 10431 10501 taagaggtaa cgctagtgaa gctaaaacac ttgcacaaca agctaaagaa actgctgctg gtttgtcaac 10571 10641 agaaattgac acagtaacat caaccgcaaa tcaagcgttg acgaaggctg gtacagcaca acaaaccgca 10711 gaacaagcga aaacaacagc aaacagtatc agcgcagttg caacggcagc taaaaacaca qctqattcaq 10781 cacaaaaaag tgcaactgat ctagctgttc gagtaagcag tttagaggac acagcaatac aatatactgt 10851 attaccatag gaggaaaaat aatggcaaat aaaaatattc aaatgaagga tagcaatgac aataatttat 10921 atccaagtgt tcgagcagaa aacttgttag atttgaccag tcgtgctgaa ttaacaatga caaattgtca 10991 attatatgca gctggtgata aaacaaatgc aatctcttat ctcggtgcag taggtatgct cgaaggtatq 11061 ataaagttta ctgaaagttt gacaaaccct gtgatcacaa cgctaccaga aggttttaga ccaataagaa caaaacgtat tggttgtttc gcaaaatatt acacaccaaa tccaacagat acaaaagaaa tggtttatgt 11131 11201 atcaatcaca cctgatggca aagtaactgt aaatgacaat gtaggtaaaa tcgaatatct atccctaqat 11271 aattgcgttt tccctctaaa ataaggaggt tcatatggaa gaacgaattg atattcaaat gaacaagatg 11341 aaagaagaaa atcaaaagaa ttacctattg caccctgaaa cgaacccgaa acaagttgtt tttgatgaaa 11411 cattgcatgg aaatgaaaat caggagagtt tcaacaattt tgttgacaca agaaaaatga caactacaat 11481 tgatgtaagt gcttatgggg ttatcgctga cggtgtaaca gattgtacac caatattaaa taaattactt 11551 gaagaaaaaa gcgaaatggg tatcactttt tattttcctc cttgtgaacg tgattcatat tatcgctttg ctaacaccat tgaattgaaa cgtgatgtac ctgtagttac tttcttagga tcgggagaaa cgacattaaa 11621 gtttgaaaca atgacggcat ttaatgtaaa catcgaaagt ttcaatattg atggttttgc attatggttg 11691 ccacaaggcg ctcaaagtgg taaaggaatt 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aagtaatgta tataactagg aggatatgag atggcaactc ttacaaatga 12671 acaaatagct agaggacaaa caatcgctaa aatactttca aaatatggct ataataaaaa ttcacaagta 12741 ggagttgteg ccaatctcca ttgggaatcg gctggtttga acccgaacag caatgaatat ggtggaggcg gatatgggtt aggtcaatgg acgcctaaaa gcaatcttta tcgccaagca caaatttgtg ggttgtctaa tgctaaagct gaaacgttgg aaggtcaagc agagatcatc gctcaagggg ataaaacagg tcaatggatg 12811 12881 gataatacac ctgtttcttc tgcaggttat actaaccctc agacctttc agcatttaaa caatctgcaa atattgatgt tgctacaatt aatttatgt gtcactggga acgccctggt aaacttcata tcgaagaaag 12951 13021 acttgatett geacaagett atagtaagea tattgaeggt ageggtggeg gtggegtaaa aegttgetat 13091 ggaaccccaa tcaagaatac aaatcttgat cctaaaagtt tcatgagtgg acaacttttt ggcacgcatg 13161 13231 caggaaacgg cagaccaaat aatttccatg atggtttgga ctttggttca attgatcacc ctggcaatga 13301 aatgattgca tgttgcgatg gaacagtaac acatgttgga acaatgggag cattaagagc gtattttgtg 13371 ataaatgatg gtacttacaa tatcgtttat caagaattta gttataacca gtcaaatata aaqqtaaaaq 13441 ttggcgacaa agttaagaac 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tcgctttatt aaacacggcg aacaatttat ttatttaaga agattcaaaa 15191 cagaacttaa aaagatteet caatttttea aaacaatgge gaaagaattt cetgateata aacttgaagt 15261 aaaaggaaaa gaattctatt gtgatgataa attaatgggt tgggctgttc cacttagtac gtggggaatt

gaaaaatcta atgaatatcc cgaagttcgt acaattttgt ttgatgagtt tttaattgag aaatcaaaaa

15401	tcacttattt	accaaacgaa	gctgaagcct	tattgaacat	gatggaaacg	gttttccgaa	gacgtacaaa
15471	tacaagatgt	gttatgttga	gtaatgcaac	tagtgtagtg	aacccttatt	tcttgtattt	caatctgcag
15541	ccagatttga	ataagcgttt	taatctatat	caagatcgag	gtatattgat	tgaattgtgt	gattcaaaag
15611	actttgcaga	agtgaagaga	gaaacacctt	ttggtagatt	gattcgtgga	acagaatacg	aagattttag
15681	tatcaacaat	gagtttgtca	atgatagtga	tacgtttatt	gaaaagagaa	gtaaaaatag	tagtttctta
15751	tgcgccattg	cttttgaagg	gaaaatcttt	gggtattgga	tagacgctga	aacaggttgt	gtctatgtga
15821	gttatgatta	tcaaccaaat	acaaatcatt	tttatgcaat	gactacgaaa	gaccatgaag	aaaatagatt
15891	gctgatgaaa	aattggcgaa	ataattatta	tctttcaaca	gtggcgaaag	cattcaagaa	tagttatctg
15961	cggtttgata	acattgttat	taagaattta	cattatgatt	tgtttaataa	gatgaaaatc	tggtaaccct
16031	attttagtag	agctaccacg	attagttcta	ttacaatgat	gaatagtaga	taacatagta	attgtagtct
16101	gcgatagttt	tgttttggtt	ctttggcgtt	agtgattttt	gctaacgcct	ttttgtttgc	ttttggatcg
16171	ggtgtgttaa	tgtagacgaa	atcttttctc	atagttcttt	ctccttatac	agttttaata	attccctgta
16241	aaatgtagct	ataggacgtc	catttctttc	tattctaacg	caattcacta	tatccatttc	taggtatata
16311	cggctatatt	ttaatgcttt	tgttaaggtg	agaggttcgg	ttttgtgtat	caaaacctcc	caaccatcta
16381	tataaaatac	tgtgatatcg	tatattggtt	ccttgtagaa	tgtagccatt	attccacctc	ctttaaatag
16451	ccttttggta	tttgtaacgc	taactgatag	cgagaaccaa	cttttacgta	tgaagttact	aatttcattg
16521	cctgacaata	cttttcaaga	atgttaaatt	gactcgattc	gggtaatagc	gttgaatgag	ttaacaaaag
16591	ttcggtgata	tttatttccg	gaacgtcgaa	atcttgtaaa	gtcccctcta	tgatctctat	tttttcattg
16661	tctgaaaggt	tacgtttaca	gtagaaacgt	aaccattcaa	ttagttcgcg	gtgttctttg	aatgttcgtg
16731	caatcatttt	aattcctcct	atttgtccgt	aatttgttta	tatccgtcat	gtttcaattg	ttccgcatag
16801	tgttcaacgc	ttttcattga	tttcgttatt	gcgatattaa	tgcaatggct	atcaagataa	acatagttat
16871	atttatcatg	tgttaacacg	aactcttttg	taacgtaatc	aatgtataaa	attaattgtt	ttcctccttg
16941	tgttatttct	gacttgatag	acgctaaact	atcgttgtca	tctttagtta	gttgatttaa	accctctaaa
17011	attaatgata	aattgttaat	catgtaaaac	actcctttta	tattaatttg	atattgatac	caccaatcga
17081	ataagattgg	tagcattgta	tcgaattaat	atgttatttc	tgtagttttc	catgaatact	cggaaataag
17151	atccatatct	aattccttta	gttcttcaaa	agataacaaa	caatattcct	catcgcctac	ctcatcaata
17221	tcaataagat	aatgtttatt	gttttcggta	tctatgatat	gataattcat	atcccactca	ttaaaggggt
17291	gaagtagaga	tacctctcct	ttttcagcta	ttaatgattt	attgttcata	tgaaacactc	cttttatatt
17361	aatttgatat	tgataccacc	aatcaaatgt	gattggtagc	attgtattaa	attaatattc	tggataattt
17431	attgagaaag	tccagttatc	atcaaatgaa	attgttttat	tttcaagtaa	ctttttagcc	tcatccacct
17501	caaattctaa	atagaggaat	ttactaagtt	tatcctcatc	tctaaaaatt	ttcatacata	ccacgttatt
17571	tgaataaatt	tctgtgtata	cgatcggttc	attcatgttt	atcatccttt	ctttattaca	tatatagtat
17641	atcatgtatt	tacatatatg	tcaatcattt	aattcattta	ttttaatgat	ttatttgatt	gtttttttat
17711	gatcctttct	ttattacatc	tatattatat	catgtatgat	tgtatttgtc	aacaattaaa	ttcatataaa
17781	tgtagtttgg	ggtcagttac	atttgtgtta	tcaaaaaaag	ataatattct	att	

Table 22

## Phage 182 ORFs list

nb	Name	Frame	Position	Size (a.a.)	Key words
1	182ORF001	2	59667780		Tail protein;
2	182ORF002	1 1	21523873		DNA polymerase;
3	182ORF003	1 1	1130512639	444	
4	182ORF004	3	46265954		Major head protein;
5	182ORF005	3	1265113700		Glycyl-Glycine endopeptidase: Lysostaphin precursor:
<u>6</u>	182ORF006	1	1499516026	343 326	Encapsidation protein; ATG/GTP-binding site motif A;  Upper collar protein;
7	182ORF007	2	77958775	292	Lysozyme; Muramidase;
8	182ORF008		1410514983		: Terminal protein;
9 10	182ORF010 182ORF009	2 2	13102155 87659601		Lower collar protein:
11	182ORF011	1-1	960710158	183	Pre-neck appendage protein;
12	182ORF012	3	1087211294	140	1 110 Floor appendage protein,
13	182ORF013	1	1045610860	134	
14	182ORF014	3	1371614108	130	Lysis protein;
15	1820RF015	2	8541225	123	Early protein;
16	182ORF018	-2	1642916737	102	
17	182ORF020	3	1015810454	98	Leucine-zipper motif;
18	1820RF019	3	43234613		Head protein;
19	1820RF016	-3	1674917033		
20	182ORF022	1	1286813149	93	
21_	1820RF023	-2	1191412189	91	
22	182ORF017	1	154426	90	
23	182ORF024	3	61746446	90	]
24	182ORF025	2	548814	88	i Early protein;
25	182ORF026	-3	1299913259	86	<u> </u>
26	182ORF027	-1	1464214896	84	l
27	182ORF028	3	1443014672	80 77	! 
28	182ORF021	-3 -1	1710617339 1619916429	76	<u> </u>
29 30	1820RF030 1820RF031	-3	83798603	74	
31	182ORF031	-3	1119511413	72	
32	182ORF033	-1	47274942	71	
33	1820RF034	-1	59516160	69	
34	182ORF029	-3	1741217606	64	
35	182ORF035	-3	1557015758	62	
36	182ORF036	-3	21272315	62	
37	1820RF037	-1	1209512280	61	
38	182ORF038	3	1476914951	60	
39	182ORF039	2	999210171	59	
40	182ORF040	-3	1602916202	57	
41	1820RF041	11	38864056	56	Early protein;
42	182ORF042	-3	1067110832	53	
43	182ORF043	-3	1049110652	53	
44	1820RF044	-1	62996457	52	
45	182ORF045	-2	65716729	52	· · · · · · · · · · · · · · · · · · ·
46	1820RF046	2	23722527	51 50	
47 48	182ORF047 182ORF048	-2 -3	1320113353 32433395	1 50 I 50	
49	182ORF049	3	15781724	48	
50	182ORF050	2	80128155	47	<del></del>
51	182ORF051	3	93909530	46	
52	182ORF052	1	40964233	45	
53	182ORF053	2	1565615793	45	
54	182ORF054	-2	80028136	44	
55	182ORF055	2	83248455	43	
56	182ORF056	3	65496680	43	
57	182ORF057	-3	81338264	43	
58	182ORF058	-1	50485176	42	· · · · · · · · · · · · · · · · · · ·
59	182ORF059	-2	1574815876	42	·
60	182ORF060	-3	1527615404	42	-
61	182ORF061	-3	19742102	42	
62 63	182ORF062	-2	18671992	41	
	182ORF063	-3	1418114306	41	

WO 00/32825

# 317

65	182ORF065	-2	34603582	1 40 1
66	182ORF066	1	42344353	39
67	1 182ORF067	-1	1376313882	39
68	182ORF068	-1	71487267	39
69	1820RF069	-3	49085027	39
70	182ORF070	-3	9121031	39
71	182ORF071	2	1174111857	38
72	182ORF072	-3	1161011723	37
73	182ORF073	-3	27632876	37
74	182ORF074	-1	88138923	36 1
75	182ORF075	-3	73537463	! 36 i
76	182ORF076	-3	23162426	36
77	182ORF077	2	1185811965	35_ :
78	182ORF078	-2	75647671	35
79	182ORF079	-2	73817488	35_ i
80	182ORF080	-2	43724473	33

- \_\_\_\_\_

### Table 23

Predicted amino acid sequences of ORFs from phage 182

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5966
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6134
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57
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6218
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85
         K W Q Y A F V T D I E Y K N D N T S F V T P E I D V L
6302
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113
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F I N T I E E S L D Y G R E Y T T N V T T F H P N D G
6386
141
       gtcaattttcttgttattctaacaagtgaagcaatgccagttggagataaggaagataaatcaggaggatcaatagtaggtggc
6470
169
       V N F L V I L T S E A M P V G D K E D K S G G S I V G G
       6554
       6638
225
       G E Y M A F L T T K E P F L N K I V G M Y V T S Y T G I
6722
       ccattcattgtggatcacgcgaacaaaacggtaaggtataatgcaggaggttcttataagatcatgcttccaacctacgctagt
253
       P F I V D H A N K.T V R Y N A G G S Y K I M L P T Y A S
6806
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281
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G N V Y N Y F R E A F P F N V K E S K L F M Y P Y C L I
6890
309
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E I T D T K G H V M T L R P E Y L T G G K L S V Y V K G
6974
337
7058
       365
7142
       {\tt aagatgttaatcgataatgatcctaacgatgtaggagttaaatctgactatgcttctgcattcatgcaaggaaacaaaaactcc}
       K M L I D N D P N D V G V K S D Y A S A F M Q G N K N S
393
7226
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       LIAQEQNIRNTFRHGMGNSAMSTGGAIF
7310
       tcagccttagcaagtaacaacccttttgttggtttgactaacatcatgggagcaggacaacaagtaaacaactatgtttctgaa
       S A L A S N N P F V G L T N I M G A G Q Q V N N Y V S E
449
7394
       477
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7478
       505
       L S F T T G N F Q N Y Y Q L R F K Q I K Y E Y A T R L D
7562
       533
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7646
       561
7730
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589
1820RF002
2152
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2320
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         H N E K F D G E F M L S W L F K N G F K W C K E A K E
2404
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85
       D R T F S T L I S N M G Q W Y A L E I C W E V N Y T T T
2488
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113
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2572
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141
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2656
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169
       E W E Y L K N D I Q I M A M A L K I Q F D Q G L T R M T
2740
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197
2824
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225
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2908
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253
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FYEGEYKPNNDYPLYIQNIKVRFRLKEG
2992
281
       3076
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309
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3160
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        D E L I D L T L T N V D L E L F F E H Y D I L E I H Y T
337
3244
        {\tt tacggatatatgttcaaagcttcttgtgatatgttcaaaggctggatcgataaatggatcgaagtaaagaacaccaccgaaggg}
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365
3328
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393
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3412
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3496
        449
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3580
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477
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3664
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D R I K E I V T F D N F E V G F S S Y G K L L P K R T Q
3748
533
3832
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1820RF003
11305
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11389
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29
11473
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57
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11557
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85
        K S E M G I T F Y F P P C E R D S Y Y R F A N T I E L K
11641
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113
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11725
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141
        ESFNIDGFALWLPQGAQSGKGIFFNDTR
11809
        169
        NYNRFDFDLFVRNCTLNEGTYVVARGR
        ggggttacatttgaaaattgtctattctctaatatctctcaagcaattatcaaaacagcttttcccgatgtaaatggtatgtgg
G V T F B N C L F S N I S Q A I I K T A F P D V N G M W
11893
197
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11977
225
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253
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281
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12229
        309
        Q D V D Q A Y I D V D V Y C R N S Q V E G M N S T A I S
12313
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337
12397
        365
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        12481
393
12565
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421
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S S S L F Q A N V P M P N A D N I E A V G A G I T R L
4710
29
4794
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57
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4878
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85
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4962
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113
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Y Y K Q T I Q E A W L E K A F T S W D N F N S F V A G V
5046
141
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M N A L Y T G D E V S E F E Y T K L L I A N Y Q E K E L
5130
169
       ttcaaagagatcgaaattggcgaaattactgaatcaaatgcaaaagaatttatccgtaagatcaaatcaacctctaacaaatta
5214
197
5298
       {\tt gaatttatgagttecgcttacaacgctcaaggagttaaaacatctacctcaaaatctgatcaatacgttat{\tt battgacgccgac}
225
       E P M S S A Y N A Q G V K T S T S K S D Q Y V I I D A D
       5382
253
5466
        gatgagtttcctaaaaaagaaggcgaagaatcgtcaaatattgtggcagttattgtagatagtgaatggtttatqatctacqac
```

```
DEFPKKEGEESSNIVAVIVDSEWFMIYD
281
5550
         aaattgtacaaaacaagtctatacaaccctgaagggttatattggaattattggttgcaccaccaccacctatattctact
309
         K L Y K T T S L Y N P E G L Y W N Y W L H H H Q L Y
        tctcaattcgggaacgctgttgcttttgttaaatcagcaacaaacctgtcacaaaagttgcttttgcaagtgcaacaactagt
S Q F G N A V A F V K S A T K P V T K V A F A S A T T S
5634
337
5718
        gttgttaaaggatcatctaaagatatcgcattgacatttacaccagtagaagcaacaaaccaacaaggagaagttgtttcatca
365
           V K G S S K D I À L T F T P V E A T N Q Q G E V V S S
        gcaccagcattggttaaggcaaccgtaaaacaacagcaggtaaagcgactgccgtaaccgtagaaggcttagaagtcggtcaa
A P A L V K A T V K Q T A G K A T A V T V E G L E V G Q
5802
393
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5886
421
         SLVTFTAIGGQQATVLVTVTSD
1820RF005
        atggcaactettacaaatgaacaaatagetagagacaaacaategetaaaataettteaaaatatggetataataaaaattea
M A T L T N E Q I A R G Q T I A K I L S K Y G Y N K N S
12651
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12735
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12819
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L G Q W T P K S N L Y R Q A Q I C G L S N A K A E T L E
57
12903
        85
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        12987
113
        PQTLSAFKQSANIDVATINFMCHWERP
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K L H I E E R L D L A Q A Y S K H I D G S G G G V K R
13071
141
13155
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        C Y G T P I K N T N L D P K S F M S G Q L F G T H A G
169
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G R P N N F H D G L D F G S I D H P G N E M I A C C D G
13239
197
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13323
        T V T H V G T M G A L R A Y F V I N D G T Y N I V Y Q E
225
13407
        F S Y N Q S N I K V K V G D K V K N G Q V C A I R D A D
253
13491
        281
        H L H L G F T K K D F M T A L G S S F I D D G T W E D P
13575
        \verb|ttgaagtttttagggcaatgttttggagatggagatactggcggagataatgacgataacaataaggataaaaatgatcttatt|
309
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13659
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337
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1820RF006
14995
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L F V I G A R G I G K T Y G Y K K F V V N R F I K H G E
15079
29
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15163
57
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        Cataaacttgaagtaaaaggaaaagaattctattgtgatgataaattaatgggttgggctgttccacttagtacgtggggaatt H K L E V K G K E F Y C D D K L M G W A V P L S T W G I
15247
85
15331
        113
        E K S N E Y P E V R T I L F D E F L I E K S K I T Y L
15415
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141
        N E A E A L L N M M E T V F R R T N T R C V M L S N A
15499
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169
               V N P Y F L Y F N L Q P D L N K R F N L Y Q D R G
15583
        197
15667
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225
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15751
253
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15835
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281
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15919
309
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16003
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337
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#### 1820RF007

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7879 agtgatgtagaagaaatcagctactatgaacattatcgtagacaactcacgctccttacgttttcagttgtttgaatgggaaaat
29 S D V E E I S Y Y E H Y R R Q L T L L T F Q L F E W E N

7963 ttgccaaaatcaattgaccctcgttatttagaaaattgcttacacactaatggttatcttggtttttaaaagaccctacactt
57 L P K S I D P R Y L E I A L H T N G Y L G F F K D P T L

8047 gggttcatggtttgcgcaggggcagaagatggtcaaatcgatcattatcacaaccctattttctttacagcaaacgaagcaatg

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8131
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113
        Y H K R Y P V L R Y D D D D D K S K C I M L Y N N D L K
8215
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        V P T L P S L H R F A L D M A D I N Q I S R V N R R A
141
8299
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253
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#### 1820RF010

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12196
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        LENGCCFGHASYAHILDNNLRHLRLCYQ
29
        aaatttcacagaaattaa 12095
12112
57
        KFHRN *
1820RF038
14769
       {\tt gtgatgagtttattttcactcttacaacaggtagcacaagcgtgttttattttgacggagaaacgatctttgaattgtctgatc}
1
        V M S L F S L L Q Q V A Q A C F I L T E K R S L N C L I
14853
        caacacaactcgatcatattagaggaacatacaatcatgttcatggaaaagaaatcccatcaatggtgtggacacctgaacaat
       Q H N S I I L E E H T I M F M E K K S H Q W
29
14937
       ttgatatttacttaa 14951
       LIFT
57
1820RF039
       9992
10076
       LLQSIKLVAGIPLQEIPILTLKERRV
29
10160
       ggcagacattag 10171
1820RF040
       16202
       MRKDFVYINTPDPKANKKALAKITNAKE
       ccaaaacaaaactatcgcagactacaattactatgttatctactattcatcattgtaatagaactaatcgtggtagctctacta
16118
       PKQNYRRLQLLCYLLFIIVIELIV
29
16034
       aaatag 16029
57
1820RF041
3886
       atggaactatataaagcaatgtttatcgtacgtgatgaaggtactattgacggttacgatactgaacactatgtagatatttct
       M E L Y K A M F I V R D E G T I D G Y D T E H Y V D I S ttacatgactttgaagaaatatatggaaaagaaacacgtgaaattgaagcagtaacattagtaaaaacacggaaatttaaaaaaa
3970
       L H D F E E I Y G K E T R E I E A V T L V K T G N L K K taa 4056
29
4054
57
1820RF042
       10832
10748
29
1820RF043
       10652
10568
1820RF044
       {\tt atgaaaagttgttacatttgttgttgttgttattctctaccgtaatcaagcgactcttcaattgtattaatgaaaggtattccatt}
6457
       M K S C Y I C C C V F S T V I K R L F N C I N E R Y S I
6373
       cgaataataaagttgagggtgttcttttgcaatgaaactttctcgtataccaatatcgaaacgataagtttgtaa 6299
       RIIKLRVFFCNETFSYTNIET
29
1820RF045
6729
       atgaatggtatacctgtatacgacgttacatacatcccgactatcttatttaaaaaaggttctttcgttgtaagaaacgccatg
       M N G I P V Y D V T Y I P T I L P K K G S F V -V R N - TA M
6645
       tactctccaaaattagcattgcctgccccatttggtttgtatacctccccacttgaattgataggaagtaaataa 6571
29
       Y S P K L A L P A P F G L Y T S P L E L I G S K
1820RF046
2372
       \verb|atggtttcaaatggtgtaaagaagcaaaagaagatcgaacattctccacactcatatcaaatatgggtcaatggtatgctttgg|
       M V S N G V K K Q K K I E H S P H S Y Q I W V N G M L W
2456
       aaatttgttgggaagttaattacacaacaacaacaacaggtaaaacgaaaaaagagaaatctcgaacaataa 2527
       K F V G K L I T Q Q Q N Q V K R K K R N L E Q .
29
```

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326
1820RF047
13353
               {\tt atgctcccattgttccaacatgtgttactgttccatcgcaacatgcaatcatttcattgccagggtgatcaattgaaccaaagtgatccattgttccattgttccattgttccattgtaccaaagtgatcaattgtaccaaagtgatccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattgttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccattttccatt
               M L P L F Q H V L L F H R N M Q S F H C Q G D Q L N Q S
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13269
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29
1820RF048
3395
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3311
               F T S I H L S I O P L N I S O E A L N I Y P
29
1820RF049
1578
               M L Q S Q E R K S K K R K L K Q S K L K K R K K N T T K agettaacaaagttgaagttaagaagcccacagaaaacacaattgtcaccaactattttaa 1724
1662
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1820RF050
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8012
                                 V S L K T L H L G S W F A Q G Q K M V K S I I I
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8096
29
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1820RF051
9390
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9474
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1820RF052
4096
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1
4180
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15740
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29
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1820RF054
8136
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              V I H C F V C C K E N R V V I M I D L T I F C P C A N H gaacccaagtgtagggtctttaaagaaaccaagataaccattagtgtgtaa 8002
8052
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1820RF055
8324
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8408
                 LTNLLMYGKQMLHM
29
1820RF056
6549
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6633
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29
              I L E S T W R F L Q R K N L F
1820RF057
8264
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              M S A I S K A K R C K L G N V G T F K S L L Y N M I H P
8180
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29
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1820RF058
5176
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gtgtattcaaattcgcttacttcgtcacctgtgtataaagcgttcattacaccagcaacgaaactattgaaattatcccatgaa V Y S N S L T S S P V Y K A F I T P A T K L L K L S H E 5092 29 V N A F S N H A S W I V C L 1820RF059 15876 M V F R S H C I K M I C I W L 1 tatccaatacccaaagattttcccttcaaaagcaatggcgcataa 15748 15792 29 Y P I P K D F P F K S N G A \* 1820RF060  $\tt gtgatttttgatttctcaattaaaaactcatcaaacaaaattgtacgaacttcgggatattcattagatttttcaattccccac$ 15404 VIFD F S I K N S S N K I V R T S G Y S L D F S I P H 15320 gtactaagtggaacagcccaacccattaatttatcatcacaatag 15276 29 V L S G T A Q P I N L S S Q \* 1820RF061 2102 atgaggggacttetecacetgttteagactegateaettttgeaatetttaetgtaaaettgttetttttetetqttqtaettetq

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1820RF062
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1992
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1
1908
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1820RF063
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14306
14222
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1820RF064
         atgatgttagtcaaaccaaccaacaaagggttgttacttgctaaggctgaaaagatcgctcctcctgtactcattgcactgtttccc M M L V K P T K G L L L A K A E K I A P P V L I A L F P ataccatgtctgaaagtattgcgaatgttttgctcttga 7234
7356
7272
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1820RF065
3582
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         M N A I C I T I N N A I K T F L S G C N G S I S T P S R
3498
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         HKTSKRNINRIS
1820RF066
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4318
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         R L W K L K N M N Q F *
29
1820RF067
13882
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         M I P A L A L K L L K S I S L L N L A I V K P N T K S I
13798
         atcattgcaaccattaaccatataatcaaaccataa 13763
29
         IIATINHIIKP *
1820RF068
7267
         atgtctgaaagtattgcgaatgttttgctcttgagcaatcaaggagtttttgtttccttgcatgaatgcagaagcatagtcaga
         M S E S I A N V L L L S N Q G V F V S L H E C R S I V R
7183
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29
         FNSYIVRIIID
1820RF069
5027
         gtggaacaatgtttttacatcgggaacttcctgtttaaatacccctgtaacagactcgtcagggttgaacttatgttcctgtgc
         V E Q C F Y I G N F L F K Y P C N R L V R V E L M F L C
1
         aatgtcaacaaaatttcttcaatcgttcgacctaa 4908
4943
         NVNKNFFNRST
29
1820RF070
1031
         {\tt gtgatggttcggctccaccaaaaccagaaacttcgtctgagtaaactagaatatctttcaattctagtacttcgccaattgcgt}
          MVRLHQNQKLRLSKLEYLSILVLRQLR
947
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29
             TELKPRLWH*
1820RF071
11741
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         MVLHYGCHKALKVVKEFSLMILAITIVL
11825
         actttgatttgtttgttcgtaactgtactttaa 11857
29
         TLICLFVTVL *
1820RF072
11723
         atgtttacattaaatgccgtcattgtttcaaactttaatgtcgtttctccccgatcctaagaaagtaactacaggtacatcacgt
         M F T L N A V I V S N F N V V S P D P K K V T T G T S R ttcaattcaatggtgttagcaaagcgataa 11610
11639
29
         FNSMVLAKR*
1820RF073
2876
         gtgaagccgcctttgtatgctttacgtaagtctttatcaaaccctaaagacaaaataggaaaccattgtttgaaagttgatttt
         V K P P L Y A L R K S L S N P K D K I G N H C L K V D F ccatgtgtagcttttagccaatctttgtaa 2763
2792
29
         PCVAFSOSL
1820RF074
8923
         gtgattgataaattttgtttcaaattctgctcgttttgtttcgtcataaaacggataatcaaaacaaatcaaacttgttttcggcc
        V I D K F C F K F C S F C F V I K R I I K I K Q L F S A aacttcaatacgttctttcgagataa 8813
8839
        NFNTFFSR *
1820RF075
        gtgttacattatotggaatattttcgatatctgccactttacctgccaagaggttcaaaccgttttcttttcagaaacatagt
7463
         V L H Y L B Y F R Y L P L Y L P R G S N R F L F Q K H S
7379
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29
        CLLVVLLP
1820RF076
2426
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M S V E N V R S S F A S L H H L K P F L N N H E S I N S
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P S N F S L W K *
2342
29
1820RF077
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11858
1
11942
29
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1820RF078
         7671
7587
         tttgtgccatacattgagaagtaa 7564
29
         F V P Y I E K *
1820RF079
         gtgaaagataagtttgatccaagctgtgttacattatctggaatattttcgatatctgccactttacctgccaagaggttcaaa V K D K F D P S C V T L S G I F S I S A T L P A K R F K ccgttttctttttcagaaacatag 7381
7488
1
7404
           FSFSET .
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1820RF080
         4473
1
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4389
         LLHRL *
```

#### Table 24

Sequence similarities phage 182 and public databases

```
Phage: 182
Database: nr
Query= sid | 110156 | lan | 1820RF001 Phage 182 ORF | 5966-7780 | 2
            (604 letters)
gi|138124|sp|P07534|VG9_BPPZA TAIL PROTEIN (LATE PROTEIN GP9) >...
                                                                                     384 e-105
gi | 138123 | sp | P04331 | VG9 BPPH2 TAIL PROTEIN (LATE PROTEIN GP9) >...
                                                                                      374
                                                                                           e-103
gi|1429238|gn1|PID|e1173412 (X99260) tail protein [Bacteriophag...
                                                                                      346 3e-94
gi|215339 (M12456) p9 tail protein (Bacteriophage phi-29) >gi|2...
                                                                                      208
                                                                                            8e-53
gi|1181970|gnl|PID|e221269 (Z47794) tail protein [Bacteriophage...
gi|1181968|gnl|PID|e221267 (Z47794) tail protein [Bacteriophage...
                                                                                            8e-09
                                                                                       56
                                                                                            6e-07
qi 2500030 sp Q59968 CARA_SULSO CARBAMOYL-PHOSPHATE SYNTHASE SM...
Query= sid | 110157 | lan | 1820RF002 Phage 182 ORF | 2152-3873 | 1
            (573 letters)
gi|118848|sp|P19894|DPOL_BPM2 DNA POLYMERASE >gi|76896|pir||JQ0...
                                                                                      665 0.0
gi|1429230|gn1|PID|e1173404 (X99260) DNA polymerase [Bacterioph...
                                                                                      657 0.0
gi|118849|sp|P03680|DPOL_BPPH2 DNA POLYMERASE (EARLY PROTEIN GP...
gi|118851|sp|P06950|DPOL_BPPZA DNA POLYMERASE (EARLY PROTEIN GP...
                                                                                      654
                                                                                            0.0
                                                                                      654 0.0
gi|15732 (X53371) DNA polymerase (AA 1-575) [Bacteriophage phi-29] gi|15734 (X53370) DNA polymerase (AA 1-575) [Bacteriophage phi-29] gi|15734 (X53370) DNA polymerase (AA 1-575) [Bacteriophage phi-29] gi|1572479|gnl|PID|e242301 (X96987) DNA polymerase [Bacteriopha... gi|1072656|pir||S51275 DNA polymerase - phage CP-1 >gi|836593|g... gi|118847|sp|P22374|DPOM_ASCIM PROBABLE DNA POLYMERASE >gi|8385... gi|461962|sp|P33537|DPOM_NEUCR PROBABLE DNA POLYMERASE >gi|2833...
                                                                                      651
                                                                                            0.0
                                                                                      651
                                                                                            0.0
                                                                                      565
                                                                                            e-160
                                                                                      301
                                                                                            1e-80
                                                                                       71
                                                                                            3e-11
                                                                                       65
                                                                                            1e-09
gi|461963|sp|P33538|DPOM_NEUIN PROBABLE DNA POLYMERASE >gi|1018...
                                                                                       62
                                                                                            1e-08
gi|1084487|pir||S41618 DNA polymerase - slime mold (Physarum po...
                                                                                       61
                                                                                            3e-08
gi|2435429 (AF012250) unassigned reading frame (possible DNA po...
                                                                                       61
                                                                                            3e-08
gi|578157|gnl|PID|e246743 (X52106) DNA polymerase [Neurospora i...
                                                                                       59
                                                                                            1e-07
gi|2147969|pir||S72369 probable DNA-polymerase - Gelasinospora ...
gi|2147968|pir||S62752 probable DNA-polymerase - Gelasinospora ...
                                                                                       58
                                                                                            2e-07
                                                                                       58
                                                                                            2e-07
gi 3511140 (AF061244) B type DNA polymerase (Agrocybe aegerita)
                                                                                       57
                                                                                            3e-07
gi | 118850 | sp | P10479 | DPOL_BPPRD DNA POLYMERASE (PROTEIN P1) >gi | ...
                                                                                       56
                                                                                            6e-07
gi|578144 (X63909) putative DNA-polymerase, B-type [Morchella c...
                                                                                            3e-04
gi|232013|sp|P30322|DPOM_AGABT_PROBABLE_DNA_POLYMERASE >gi|3208...
                                                                                            6e-04
Query= sid|110159|lan|182ORF004 Phage 182 ORF|4626-5954|3
           (442 letters)
gi|138117|sp|P13849|VG8_BPPH2 MAJOR HEAD PROTEIN (LATE PROTEIN ...
                                                                                      309
                                                                                            2e-83
gi|138118|sp|P07531|VG8_BPPZA MAJOR HEAD PROTEIN (LATE PROTEIN ...
                                                                                     305
                                                                                            3e-82
gi|1429236|gn1|PID|e1173410 (X99260) major head protein [Bacter...
                                                                                     300
                                                                                            1e-80
gi|1181958|gnl|PID|e221257 (247794) major head protein [Bacteri...
                                                                                     152
                                                                                            6e-36
Query= sid|110160|lan|1820RF005 Phage 182 ORF|12651-13700|3
           (349 letters)
gi|137932|sp|P15132|VG13_BPPH2 MORPHOGENESIS PROTEIN 1 (LATE PR...
                                                                                            8e-06
gi|1429242|gnl|PID|e1173416 (X99260) morphogenesis protein [Bac...
                                                                                       48
                                                                                            7e-05
gi|137933|sp|P07538|VG13_BPPZA MORPHOGENESIS PROTEIN 1 (LATE PR...
                                                                                       47 2e-04
Query= sid|110161|1an|182ORF006 Phage 182 ORF|14995-16026|1
           (343 letters)
qi|137944|sp|P11014|VG16 BPPH2 ENCAPSIDATION PROTEIN (LATE PROT...
                                                                                     402 e-111
gi 137945 sp P07541 VG16 BPPZA ENCAPSIDATION PROTEIN (LATE PROT...
                                                                                     402 e-111
gi|1429245|gn1|PID|e1173419 (X99260) encapsidation protein [Bac...
                                                                                     381
                                                                                            e-105
gi | 1181972 | gn1 | PID | e221271 (Z47794) encapsidation protein [Bact...
                                                                                     159 2e-38
```

(123 letters)

```
gi|1429239|qn1|PID|e1173413 (X99260) upper collar protein [Bact...
                                                                                 271 5e-72
gi|137915|sp|P07535|VG10 BPPZA UPPER COLLAR PROTEIN (CONNECTOR ... gi|137914|sp|P04332|VG10_BPPH2 UPPER COLLAR PROTEIN (CONNECTOR ...
                                                                                 256 le-67
                                                                                 256 2e-67
gi|1181960|gn1|PID|e221259 (Z47794) connector protein [Bacterio...
                                                                                 148
                                                                                      6e-35
 Query= sid|110163|lan|182ORF008 Phage 182 ORF|14105-14983|2
           (292 letters)
gi|4210750|gn1|PID|e1374037 (AJ132604) LysL protein (Lactococcu...
                                                                                 139
                                                                                      2e-32
gi 462559 sp P34020 LYC_CLOAB AUTOLYTIC LYSOZYME (1,4-BETA-N-AC...
                                                                                  75
                                                                                      8e-13
gi|2327014 (U82823) putative lysozyme (Saccharopolyspora erythr...
gi|126652|sp|P25310|LYCM_STRGL LYSOZYME M1 PRECURSOR (1,4-BETA-...
                                                                                      2e-09
                                                                                  60
                                                                                      2e-08
gi 127789 sp P19386 LYCA BPCP9 LYSOZYME (ENDOLYSIN) (MURAMIDASE...
                                                                                  60
                                                                                      2e-08
gi 67761 pir | MUBPCP N-acetylmuramoyl-L-alanine amidase (EC 3.5...
                                                                                  59
                                                                                      3e-08
qi|4105636 (AF049087) lys (Leuconostoc oenos bacteriophage 10MC)
                                                                                  59
                                                                                      3e-08
gi|623084 (L02496) muramidase; muramidase [Bacteriophage LL-H]
                                                                                  57
                                                                                      1e-07
gi | 127787 | sp | P15057 | LYCA BPCP1 LYSOZYME (ENDOLYSIN) (MURAMIDASE...
                                                                                  57
                                                                                      2e-07
gi 126597 sp P00721 LYCH CHASP N, O-DIACETYLMURAMIDASE (LYSOZYME...
                                                                                  57
                                                                                      26-07
gi 127788 sp P19385 LYCA_BPCP7 LYSOZYME (ENDOLYSIN) (MURAMIDASE...
                                                                                  57
                                                                                      2e-07
gi 67762 pir | MUBPC7 N-acetylmuramoyl-L-alanine amidase (EC 3.5...
                                                                                  56
                                                                                      3e-07
gi 3025168 sp P76421 YEGK_ECOLI HYPOTHETICAL 32.0 KD PROTEIN IN...
                                                                                  53
                                                                                      2e-06
gi 4204413 (AF047001) Lys44 [Oenococcus oeni temperate bacterio...
                                                                                  53
                                                                                      3e-06
gi 2116978 | gnl | PID | d1020940 (D88151) cortical fragment-lytic en...
                                                                                  52
                                                                                      5e-06
gi|2392844 (AF011378) lysin [Bacteriophage sk1]
                                                                                  48
                                                                                      8e-05
Query= sid|110164|lan|182ORF009 Phage 182 ORF|8765-9601|2
           (278 letters)
gi|1429240|gnl|PID|e1173414 (X99260) lower collar protein (Bact...
                                                                                180 le-44
gi 137921 sp P04333 VG11_BPPH2 LOWER COLLAR PROTEIN (LATE PROTE...
                                                                                171
                                                                                      5e-42
gi|215341 (M12456) pl1 lower collar protein [Bacteriophage phi-29]
                                                                                 98
                                                                                      9e-20
gi|224162|prf||1011232B protein pl1,lower collar {Bacteriophage...
                                                                                      le-19
gi|535260 (Z30339) STARP antigen [Plasmodium reichenowi]
gi|4049753 (AF063866) ORF MSV230 hypothetical protein [Melanopl...
                                                                                      1e-05
                                                                                      4e-05
gi|2131557|pir||S70306 hypothetical protein YEL077c - yeast (Sa...
gi|131782|sp|P12753|RA50_YEAST DNA REPAIR PROTEIN RAD50 (153 KD...
                                                                                      5e-05
                                                                                 48
                                                                                      7e-05
gi|2131309|pir||S70305 hypothetical protein YBL113c - yeast (Sa...
                                                                                 47
                                                                                      2e-04
gi|499325 (Z26314) STARP antigen [Plasmodium falciparum]
gi|3845171 (AE001391) ribosome releasing factor (OO, TP) [Plasm...
gi|731903|sp|P40434|YIR7_YEAST HYPOTHETICAL 197.5 KD PROTEIN IN...
gi|1632829|gnl|PID|e276379 (Y08924) AARP2 protein [Plasmodium f...
                                                                                 46
                                                                                      3e-04
                                                                                 46
                                                                                      3e-04
                                                                                 45
                                                                                      5e-04
                                                                                 45
                                                                                      5e-04
gi 1176490 sp P40889 YJW5 YEAST HYPOTHETICAL 197.6 KD PROTEIN I...
                                                                                      5e-04
                                                                                 45
gi | 1077300 | pir | | S51848 hypothetical protein HRD1054 - yeast (Sa...
                                                                                 45
                                                                                      5e-04
gi|2425143 (AF020407) WimA [Dictyostelium discoideum]
gi|1181961|gnl|PID|e221260 (Z47794) collar protein [Bacteriopha...
                                                                                 45
                                                                                      6e-04
                                                                                 45
                                                                                      6e-04
gi|2132657|pir||S64819 probable membrane protein YLL067c - yeas...
                                                                                 45
                                                                                      8e-04
gi 2133041 pir | S65341 probable membrane protein YPR204w - yeas...
                                                                                 45
                                                                                      8e-04
gi 730275 sp P39793 PBPA BACSU PENICILLIN-BINDING PROTEINS 1A/1...
                                                                                      8e-04
Query= sid|110165|lan|1820RF010 Phage 182 ORF|1310-2155|2
           (281 letters)
gi|135604|sp|P06812|TERM_BPNF DNA TERMINAL PROTEIN >gi|75815|pi...
                                                                                 69 3e-11
gi|1572478|gn1|PID|e242334 (X96987) terminal protein (Bacteriop...
                                                                                 65
                                                                                      3e-10
gi|1429231|gn1|PID|e1173405 (X99260) terminal protein [Bacterio...
                                                                                      1e-09
Query= sid|110166|lan|1820RF011 Phage 182 ORF|9607-10158|1
          (183 letters)
gi|137928|sp|P07537|VG12 BPPZA PRE-NECK APPENDAGE PROTEIN (LATE...
gi|1429241|gnl|PID|e1173415 (X99260) pre-neck appendage protein...
gi|137927|sp|P20345|VG12_BPPH2 PRE-NECK APPENDAGE PROTEIN (LATE...
                                                                                 50 le-05
Query= sid|110169|lan|1820RF014 Phage 182 ORF|13716-14108|3
          (130 letters)
                                                                                                ____
gi|137936|ap|P11188|VG14_BPPH2 LYSIS PROTEIN (LATE PROTEIN GP14...
                                                                                 97 6e-20
gi 137938 sp P07539 VG14 BPPZA LYSIS PROTEIN (LATE PROTEIN GP14...
                                                                                     Be-20 →
                                                                                 96
gi|1429243|gn1|PID|e1173417 (X99260) lysis protein [Bacteriopha...
                                                                                 96
                                                                                     Be-20
gi 215332 (M14782) lysis protein [Bacteriophage phi-29]
                                                                                     5e-19
Query= sid | 110170 | lan | 1820RF015 Phage 182 ORF | 854-1225 | 2
```

gi 15670 (V01155) reading frame 10 (may be gene 4) [Bacteriopha gi 138072 sp P06953 VG5A_BPPZA EARLY PROTEIN GP5A >gi 75836 pir		5e-12 7e-12
Query= sid 110174 lan 1820RF019 Phage 182 ORF 4323-4613 3 (96 letters)		
gi 1429235 gnl PID e1173409 (X99260) head morphogenesis protein gi 138111 sp P13848 VG7_BPPH2 HEAD MORPHOGENESIS PROTEIN (LATE gi 138112 sp P07533 VG7_BPPZA HEAD MORPHOGENESIS PROTEIN (LATE Query= sid 110180 lan 1820RF025 Phage 182 ORF 548-814 2 (88 letters)	57	2e-09 3e-08 1e-07
gi 138099 sp P06955 VG6_BPPZA EARLY PROTEIN GP6 >gi 75841 pir   gi 138098 sp P03685 VG6_BPPH2 EARLY PROTEIN GP6 >gi 75840 pir   gi 1429234 gn1 PID e1173408 (X99260) gene 6 product [Bacterioph	54	7e-08 2e-07 2e-07

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# Table 25

# Homologies between 182 ORFs and proteins in public databases

Phage: 182			
Database: Swissprot			
Query= sid 110156 1an 1820RF001 Phage 182 ORF 5966-7780 2 (604 letters)			
gi 138124 sp P07534 VG9_BPPZA TAIL PROTEIN (LATE PROTEIN GP9) gi 138123 sp P04331 VG9_BPPH2 TAIL PROTEIN (LATE PROTEIN GP9) gi 2500030 sp Q59968 CARA_SULSO CARBAMOYL-PHOSPHATE SYNTHASE SM	374	e-106 e-103 2e-05	
Query= sid 110157 lan 1820RF002 Phage 182 ORF 2152-3873 1 (573 letters)			
gi 118848 sp P19894 DPOL_BPM2 DNA POLYMERASE gi 118849 sp P03680 DPOL_BPPH2 DNA POLYMERASE (EARLY PROTEIN GP2) gi 118851 sp P06950 DPOL_BPPZA DNA POLYMERASE (EARLY PROTEIN GP2) gi 118851 sp P06950 DPOL_BPPZA DNA POLYMERASE (EARLY PROTEIN GP2) gi 118847 sp P23374 DPOM_ASCIM PROBABLE DNA POLYMERASE gi 461962 sp P33537 DPOM_NEUCR PROBABLE DNA POLYMERASE gi 461963 sp P33538 DPOM_NEUIN PROBABLE DNA POLYMERASE gi 118850 sp P10479 DPOL_BPPRD DNA POLYMERASE (PROTEIN P1) gi 232013 sp P30322 DPOM_AGABT PROBABLE DNA POLYMERASE gi 118887 sp P10582 DPOM_MAIZE DNA POLYMERASE (S-1 DNA ORF 3)	654 654 71 65 62 56 46	0.0 0.0 0.0 7e-12 3e-10 3e-09 2e-07 2e-04 2e-04	
Query= sid 110159 lan 182ORF004 Phage 182 ORF 4626-5954 3 (442 letters)			
gi 138117 sp P13849 VG8_BPPH2 MAJOR HEAD PROTEIN (LATE PROTEIN gi 138118 sp P07531 VG8_BPPZA MAJOR HEAD PROTEIN (LATE PROTEIN			
Query= sid 110160 lan 182ORF005 Phage 182 ORF 12651-13700 3 (349 letters)			
gi 137932 sp P15132 VG13_BPPH2 MORPHOGENESIS PROTEIN 1 (LATE PR gi 137933 sp P07538 VG13_BPPZA MORPHOGENESIS PROTEIN 1 (LATE PR			
Query= sid 110161 1an 182ORF006 Phage 182 ORF 14995-16026 1 (343 letters)			
gi 137945 sp P07541 VG16_BPPZA ENCAPSIDATION PROTEIN (LATE PROT gi 137944 sp P11014 VG16_BPPH2 ENCAPSIDATION PROTEIN (LATE PROT			
Query= sid 110162 1an 1820RF007 Phage 182 ORF 7795-8775 1 (326 letters)			
gi 137915 sp P07535 VG10_BPPZA UPPER COLLAR PROTEIN (CONNECTOR gi 137914 sp P04332 VG10_BPPH2 UPPER COLLAR PROTEIN (CONNECTOR		3e-68 5e-68	
Query= sid 110163 1an 182ORF008 Phage 182 ORF 14105-14983 2 (292 letters)			
gi 462559 sp P34020 LYC_CLOAB AUTOLYTIC LYSOZYME (1,4-BETA-N-AC gi 126652 sp P25310 LYCM_STRGL LYSOZYME M1 PRECURSOR (1,4-BETA gi 127789 sp P19386 LYCA_BPCP9 LYSOZYME (ENDOLYSIN) (MURAMIDASE gi 127787 sp P15057 LYCA_BPCP1 LYSOZYME (ENDOLYSIN) (MURAMIDASE gi 126597 sp P00721 LYCH_CHASP N,O-DIACETYLMURAMIDASE (LYSOZYME gi 127788 sp P19385 LYCA_BPCP7 LYSOZYME (ENDOLYSIN) (MURAMIDASE gi 3025168 sp P76421 YEGX_ECOLI HYPOTHETICAL 32.0 KD PROTEIN IN	60 57 57 57	5e-09 5e-09 4e-08	
Query= sid 110164 lan 182ORF009 Phage 182 ORF 8765-9601 2 (278 letters)		-	
gi 137921 sp P04333 VG11_BPPH2 LOWER COLLAR PROTEIN (LATE PROTE gi 131782 sp P12753 RA50_YEAST DNA REPAIR PROTEIN RAD50 (153 KD gi 1176490 sp P40889 YJW5_YEAST HYPOTHETICAL 197.6 KD PROTEIN I gi 731903 sp P40434 YIR7_YEAST HYPOTHETICAL 197.5 KD PROTEIN IN gi 730275 sp P39793 PBPA_BACSU PENICILLIN-BINDING PROTEINS 1A/1	171 48 45 45	le-04 2e-04	
gi 1168610 sp P41696 AZF1_YEAST ASPARAGINE-RICH ZINC FINGER PRO	44	3e-04	

333		
gi 731587 sp P38900 YH19_YEAST HYPOTHETICAL 70.1 KD PROTEIN IN	44	3e-04
Query= sid 110165 lan 182ORF010 Phage 182 ORF 1310-2155 2 (281 letters)		
gi 135604 ap P06812 TERM_BPNF DNA TERMINAL PROTEIN	69	8e-12
Query= sid 110166 lan 1820RF011 Phage 182 ORF 9607-10158 1 (183 letters)		
gi 137928 sp P07537 VG12_BPPZA PRE-NECK APPENDAGE PROTEIN (LATE gi 137927 sp P20345 VG12_BPPH2 PRE-NECK APPENDAGE PROTEIN (LATE		2e-06 3e-06
Query= sid 110169 lan 1820RF014 Phage 182 ORF 13716-14108 3 (130 letters)		
gi 137936 sp P11188 VG14_BPPH2 LYSIS PROTEIN (LATE PROTEIN GP14)	97	2e-20
gi 137938 sp P07539 VG14_BPPZA LYSIS PROTEIN (LATE PROTEIN GP14)	96	2e-20
Query= sid 110170 lan 1820RF015 Phage 182 ORF 854-1225 2 (123 letters)		
gi 138072 ap P06953 VG5A_BPPZA EARLY PROTEIN GP5A	69	2e-12
Query= sid 110174 lan 1820RF019 Phage 182 ORF 4323-4613 3 (96 letters)		
gi 138111 sp P13848 VG7_BPPH2 HEAD MORPHOGENESIS PROTEIN (LATE gi 138112 sp P07533 VG7_BPPZA HEAD MORPHOGENESIS PROTEIN (LATE	57 54	
Query= sid 110180 lan 182ORF025 Phage 182 ORF 548-814 2 (88 letters)		
gi 138099 sp P06955 VG6_BPPZA EARLY PROTEIN GP6 gi 138098 sp P03685 VG6_BPPH2 EARLY PROTEIN GP6	55 54	2e-08 5e-08

- \_\_\_\_\_

334

BLASTP 2.0.8 [Jan-05-1999]

Query= sid|110156|lan|1820RF001 Phage 182 ORF|5966-7780|2 (604 letters)

>gi|138124|sp|P07534|VG9\_BPPZA TAIL PROTEIN (LATE PROTEIN GP9) >gi|75849|pir||WMBP9Z gene 9 protein - phage PZA >gi|216058 (M11813) tail protein [Bacteriophage PZA] Length = 599

Score = 384 bits (975), Expect = e-105 Identities = 231/610 (37%), Positives = 344/610 (55%), Gaps = 36/610 (5%)

Query: 6 TNVKLLANVPFDNTYTHTRWFKTQQEQESYFNSFPVLNENRDCSYQRDTQLGGVFRVDKH 65 TNV++LA+VPF N Y +TRWF + Q ++FNS + E ++Q + V Sbjct: 9 TNVRILADVPFSNDYKNTRWFTSSSNQYNWFNSKTRVYEMSKVTFQGFRENKSYISVSLR 68

Query: 66 KDALYACNYLIFKNEETYPSKWQYAFVTDIEYKNDNTSFVTFEIDVLQTYRFDIGIRESF 125 D LY +Y++F+N + Y +KW YAFVT++EYKN T++V FEIDVLQT+ F+I +ESF

Sbjct: 69 LDLLYNASYIMFQNAD-YGNKWFYAFVTELEYKNVGTTYVHFEIDVLQTWMFNIKFQESF 127

Query: 126 IAKEHPQLYYSNGIPFINTIEESLDYGREYTTTNVTTFHPNDGVNFLVILTSEAM--PVG 183 I +EH +L+ +G P INTI+E L+YG EY +V P D + FLV+++ M G Sbjct: 128 IVREHVKLWNDDGTPTINTIDEGLNYGSEYDIVSVENHRPYDDMMFLVVISKSIMHGTAG 187

Query: 184 DKEDKSG---GSIVGGPSPFSYYLLPINSSGEVYKPN-GAGNANFGEYMAFLT---TKEP 236 S+ G P P YY+ P G+V K G NAN + LT

Sbjct: 188 EAESRLNDINASLNGMPOPLCYYIHPFYKDGKVPKTFIGDNNANLSPIVNMLTNIFSOKS 247

Query: 237 FLNKIVGMYVTSYTGIPFIVDHANKTVRYNAGGSYKIMLPTYASDPTGTMKTFAFFCVKE 296 +N IV MYVT Y G+ + +K ++ + + A D G + T

Sbjct: 248 AVNNIVNMYVTDYIGLKLDYKNGDKELKLDKDMFEQAGI---ADDKHGNVDTIF---VKK 301

Query: 297 ARTFVPKRIDLVGNVYNYFREAFPFNVKESKLFMYPYCLIEITDTKGHVMTLRPEYLTGG 356

+ ID G+ + F + +ESKL MYPYC+ E+TD KG+ M L+ EY+
Sbjct: 302 IPDYETLEID-TGDKWGGFTKD-----QESKLMMYPYCVTEVTDFKGNHMNLKTEYIDNN 355

Query: 357 KLSVYVKGSLGISNKVMIEPIDYDVSNSTI----ITNLSDKMLIDNDPNDVGVKSDYASA 412 KL + V+GSLG+SNKV DY+ S +T D LI+N+PND+ + +DY SA

Sbjct: 356 KLKIQVRGSLGVSNKVAYSIQDYNAGGSLSGGDRLTASLDTSLINNNPNDIAIINDYLSA 415

Query: 413 FMQGNKNSLIAQEQNIRNTFRHGMGNSAMSTGGAIFSALASNNPFVGLTNIMGAGQQVNN 472

++QGNKNSL Q+ +I GM +S G ++ +PF +++ G N
Sbjct: 416 YLQGNKNSLENQKSSILFNGIVGMLGGGVSAG----ASAVGRSPFGLASSVTGMTSTAGN 471

Query: 473 YVSEKENGLNLLAGKVADIENIPDNVTQLGSNLSFTTGN-FQNYYQLRFKQIKYEYATRL 531

V + + L K ADI NIP +T++G N +F GN ++ Y ++ KQ+K EY L
Sbjct: 472 AVLD----MQALQAKQADIANIPPQLTKMGGNTAFDYGNGYRGVYVIK-KQLKAEYRRSL 526

Query: 532 DRYFSMYGTKSNRVATPNLQTRKAWNFIKLKEPNIVGTMSNDVLTRVKQIFSAGVTLWHT 591

+F YG K NRV PNL+TRKA+N+I+ K+ I G ++N+ L ++ IF G+TLWHT
Sbjct: 527 SSFFHKYGYKINRVKKPNLRTRKAYNYIQTKDCFISGDINNNDLQEIRTIFDNGITLWHT 586

Query: 592 NDVLNYNQDN 601

+D+ NY+ +N

Sbjct: 587 DDIGNYSVEN 596

Query= sid|110157|lan|1820RF002 Phage 182 ORF|2152-3873|1

>gi|118848|sp|P19894|DPOL\_BPM2 DNA POLYMERASE >gi|76896|pir||JQ0161 DNA-directed DNA polymerase (EC 2.7.7.7) - phage M2 >gi|215509 (M33144) DNA polymerase (Bacteriophage M2) Length = 572

Score = 665 bits (1697), Expect = 0.0 Identities = 327/589 (55%), Positives = 420/589 (70%), Gaps = 38/589 (6%)

KKYTGDFETTTDLNDCRVWSWGVCDIDNVDNMTFGLEIDSFFEWCKMOGSTDIYFHNEKF 62 K ++ DFETTT L+DCRVW++G +I N+DN G +D F +W M+ D+YFHN KF

Sbjct: 4

```
KMFSCDFETTTKLDDCRVWAYGYMEIGNLDNYKIGNSLDEFMQWV-MEIQADLYFHNLKF 62
Query: 63 DGEFMLSWLFKNGFKWCKEAKEDRTFSTLISNMGQWYALEICWEVNYXXXXXXXXXXXXX 122
                                  + T++T+IS MGQWY ++IC+
           DG F+++WL ++GFKW E
Sbjct: 63 DGAFIVNWLEQHGFKWSNEGLPN-TYNTIISKMGQWYMIDICFGYK------GKRKL 112
Query: 123 XXIIYDSLKKYPFPVKQIAEAFNFPIKKGEIDYTKERPIGYKPTKDEWEYLKNDIQIMAM 182
+IYDSLKK PFPVK+IA+ F P+ KG+IDY ERP+G++ T +E+EY+KNDI+I+A
Sbjct: 113 HTVIYDSLKKLPFPVKKIAKDFQLPLLKGDIDYHTERPVGHEITPEEYEYIKNDIBIIAR 172
Query: 183 ALKIQFDQGLTRMTRGSDALGDYKDWLKATHGKSTFKQWFPILSLGFDKDLRKAYKGGFT 242
            AL IOF OGL RMT GSD+L +KD L
                                                F + FP LSL DK++RKAY+GGFT
Sbjct: 173 ALDIQFKQGLDRMTAGSDSLKGFKDILST----KKFNKVFPKLSLPMDKEIRKAYRGGFT 228
Query: 243 WVNKVFQGKEIGDGIVFDVNSLYPSQMYVRPLPYGTPLFYEGEYKPNNDYPLYIQNIKVR 302
            W+N ++ KEIG+G+VFDVNSLYPSQMY RPLPYG P+ ++G+Y+ + YPLYIQ I+
Sbjct: 229 WLNDKYKEKEIGEGMVFDVNSLYPSQMYSRPLPYGAPIVFQGKYEKDEQYPLYIQRIRFE 288
Query: 303 FRLKEGYIPTIQVKQSSLFIQNEYLESSVNKLGVDELIDLTLTNVDLELFFEHYDILEIH 362
            F LKEGYIPTIQ+K++ F NEYL++S GV E ++L LTNVDLEL EHY++
Sbjct: 289 FELKEGYIPTIQIKKNPFFKGNEYLKNS----GV-EPVELYLTNVDLELIQEHYELYNVE 343
Query: 363 YTYGYMFKASCDMFKGWIDKWIEVKNTTEGARKANAKGMLNSLYGKFGTNPDITGKVPYM 422
            Y G+ F+ +FK +IDKW VK EGA+K AK MLNSLYGKF +NPD+TGKVPY+
Sbjct: 344 YIDGFKFREKTGLFKDFIDKWTYVKTHEEGAKKQLAKLMLNSLYGKFASNPDVTGKVPYL 403
Query: 423 GEDGIVRLTLGEEELRDPVYVPLASFVTAWGRYTTITTAQKCFDRIIYCDTDSIHLVGTE 482
             +DG + +G+EE +DPVY P+ F+TAW R+TTIT AQ C+DRIIYCDTDSIHL GTE
Sbjct: 404 KDDGSLGFRVGDEEYKDPVYTPMGVFITAWARFTTITAAQACYDRIIYCDTDSIHLTGTE 463
Query: 483 VPEAIDHLVDPKKLGYWGHESTFQRAKFIRQKT----YVEEIDGEL----- 524
            VPE I +VDPKKLGYW HESTF+RAK++RQKT
Sbjct: 464 VPEIIKDIVDPKKLGYWAHESTFKRAKYLRQKTYIQDIYVKEVDGKLKECSPDEATTTKF 523
Query: 525 NVKCAGMPDRIKEIVTFDNFEVGFSSYGKLLPKRTQGGVVLVDTMFTIK 573
+VKCAGM D IK+ VTFDNF VGFSS GK P + GGVVLVD++FTIK
Sbjct: 524 SVKCAGMTDTIKKKVTFDNFAVGFSSMGKPKPVQVNGGVVLVDSVFTIK 572
Query= sid|110159|lan|1820RF004 Phage 182 ORF|4626-5954|3
          (442 letters)
>gi|138117|sp|P13849|VG8_BPPH2 MAJOR HEAD PROTEIN (LATE PROTEIN GP8)
            >gi|75845|pir||WMBP89 gene 8 protein - phage phi-29
           9gi|215325 (M14782) major head protein [Bacteriophage
phi-29] >gi|225362|prf||1301270B gene 8 [Bacillus sp.]
           Length = 448
 Score = 309 bits (783), Expect = 2e-83
 Identities = 176/440 (40%), Positives = 250/440 (56%), Gaps = 27/440 (6%)
           KITEQDVLRATNVETPVQLMTAIYNSSSSLFQANVPMPNADNIEAVGAGITRLDVVKNEF 63 +IT DV + + ++ AI NS F++ VP+ A+N+ VGAGI V+N+F
Query: 4
           RITFNDVKTSLGITESYDIVNAIRNSQGDNFKSYVPLATANNVAEVGAGILINQTVQNDF 61
Sbict: 2
Query: 64 ISTLVDRIGKVVIRYKSWRNPLKMFKKGNMPLGRTIEEIFVDIAQEHKFNPDESVTGVFK 123
           I++LVDRIG VVIR S NPLK FKKG +PLGRTIEEI+ DI +E +++ +E+
Sbjct: 62 ITSLVDRIGLVVIRQVSLNNPLKKFKKGQIPLGRTIEEIYTDITKEKQYDAEEAEQKVFE 121
Query: 124 QEVPDVKTLFHEINREGYYKQTIQEAWLEKAFTSWDNFNSFVAGVMNALYTGDEVSEFEY 183
+E+P+VKTLFHE NR+G+Y QTIQ+ L+ AF SW NF SFV+ ++NA+Y EV E+EY
Sbjct: 122 REMPNVKTLFHERNRQGFYHQTIQDDSLKTAFVSWGNFESFVSSIINAIYNSAEVDEYEY 181
Query: 184 TKLLIANYQEKELFKEIEIGEITESNA--KEFIRKIKSTSNKLEFM--SSAYNAQGVKTS 239
KLL+ NY K LF ++I E T S EF++K+++T+ KL S +N+ V+T
Sbjct: 182 MKLLVDNYYSKGLFTTVKIDEPTSSTGALTEFVKKMRATARKLTLPQGSRDWNSMAVRTR 241
FNM++TDF+G+ VID F
Sbjct: 242 SYMEDLHLIIDADLEAELDVDVLAKAFNMNRTDFLGNVTVIDGF-----ASTGLEAVLV 295
Query: 300 DSEWFMIYDKLYKTTSLYNPEGLYWNYWLHHHQLYSTSQFGNAVAFVKSATKPVTKVAFA 359
           D +WFM+YD L+K ++ NP GLYWNY+ H Q S S+P NAVAFV
                                                                   VT+V
Sbjct: 296 DKDWFMVYDNLHKMETVRNPRGLYWNYYYHVWQTLSVSRFANAVAFVSGDVPAVTQVIVS 355
Query: 360 SATTSVVKGSSKDIALTFTPVEATNQQGEVVSSAPALVKATVKQTAGKATAVTVEGLEVG 419
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V ATN + V
                                                        V G +T +
Sbjct: 356 PNIAAVKQGGQQQFT---AYVRATNAKDHKV--------WSVEGGSTGTAI----TG 398
Query: 420 QSLVTFTAIGGQQATVLVTV 439
                        Q TV TV
              L++ +
Sbjct: 399 DGLLSVSGNEDNQLTVKATV 418
Query= sid|110160|lan|182ORF005 Phage 182 ORF|12651-13700|3
          (349 letters)
>gi|137932|sp|P15132|VG13_BPPH2 MORPHOGENESIS PROTEIN 1 (LATE
            PROTEIN GP13) >gi | 75858 |pir | WMBP23 gene 13 protein
            phage phi-29 >gi|215331 (M14782) morphogenesis protein
            [Bacteriophage phi-29] >gi|225368|prf||1301270H gene 13
            [Bacteriophage phi-29]
            Length = 365
 Score = 51.5 bits (121), Expect = 8e-06
 Identities = 44/166 (26%), Positives = 70/166 (41%), Gaps = 14/166 (8%)
Query: 6 NEQIARGQTIAKILSKYGYNKNSQVGVVANLHWESA---GLNPNSNEXXXXXXXXXXQWT 61
+E Q I LS G+ K + G++ N+ ES GL N +E QWT
Sbjct: 12 SEMKVNAQYILNYLSSNGWTKQAICGMLGNMQSESTINPGLWQNLDEGNTSLGFGLVQWT 71
Query: 62 PKSNLYRQAQICGLSNAKAETLEGQAEIIAQGDKTGQWMDNTPVSSAGYTNPQTLSAFKQ 121 P SN A GL ++ II + + QW++ ++ Y K Sbjct: 72 PASNYINWANSQGLPYKDMDS--ELKRIIWEVNNNAQWINLRDMTFKEY------IKS 121
Query: 122 SANIDVATINFMCHWERPGKLHIEERLDLAQAYSKHIDGSGGGGVK 167
                    + F+ +ERP + ER D A+ + K++ G GGGG++
Sbjct: 122 TKTPRELAMIFLASYERPANPNQPERGDQAEYWYKNLSGGGGGGLQ 167
Query= sid | 110161 | lan | 1820RF006 Phage 182 ORF | 14995-16026 | 1
          (343 letters)
>gi|137945|sp|P07541|VG16 BPPZA ENCAPSIDATION PROTEIN (LATE PROTEIN
            GP16) >gi|75861|pir||WMBP16 gene 16 protein - phage PZA
            >gi|216065 (M11813) morphogenesis protein C
            [Bacteriophage PZA]
            Length = 332
 Score = 402 bits (1023), Expect = e-111
 Identities = 186/332 (56%), Positives = 244/332 (73%), Gaps = 2/332 (0%)
Query: 11 EKNLYYNPNNALGFNCLMLFVIGARGIGKTYGYKKFVVNRFIKHGEQFIYLRRFKTELKK 70
            +K+L+YNP L ++ ++ FVIGARGIGK+Y K + +NRFIK+GEQFIY+RR+K EL K
            DKSLFYNPQKMLSYDRILNFVIGARGIGKSYAMKVYPINRFIKYGEQFIYVRRYKPELAK 61
Query: 71 IPQFFKTMAKEFPDHKLEVKGKEFYCDDKLMGWAVPLSTWGIEKSNEYPEVRTILFDEFL 130
              +F +A+EFPDH+L VKG+ FY D. KL GWA+PLS W EKSN YP V TI+FDEF+
Sbjct: 62 VSNYFNDVAQEFPDHELVVKGRRFYIDGKLAGWAIPLSVWQSEKSNAYPNVSTIVFDEFI 121
Query: 131 IEKSKITYLPNEAEALLNMMETVFRRRTNTRCVMLSNATSVVNPYFLYFNLQPDLNKRPN 190
                  Y+PNE ALIN+M+TVFR R RC+ LSNA SVVNPYFL+FNL PD+NKPFN
Sbjct: 122 REKDNSNYIPNEVSALLNLMDTVFRNRERVRCICLSNAVSVVNPYFLFFNLVPDVNKRFN 181
Query: 191 LYQDRGILIELCDSKDFAEVKRETPFGRLIRGTEYEDFSINNEFVNDSDTFIEKRSKNSS 250
            +Y D LIE+ DS DF+ +R+T FGRLI GTEY + S++N+F+ DS FIEKRSK+S
Sbjct: 182 VYDD--ALIEIPDSLDFSSERRKTRFGRLIDGTEYGEMSLDNQFIGDSHVFIEKRSKDSK 239
Query: 251 FLCAIAFEGKIFGYWIDAETGCVYVSYDYQPNTNHFYAMTTKDHEENRLLMKNWRNNYYL 310
            F+ +I + G G W+D G +YV + P+T + Y +TT D EN +L+ N++NNY+L
Sbjct: 240 FVFSIVYNGFTLGVWVDVNQGLMYVDTAHDPSTKNVYTLTTDDLNENMMLITNYKNNYHL 299
Query: 311 STVAKAPKNSYLRFDNIVIKNLHYDLFNKMKI 342
              +A AF N YLRFDN VI+N+ Y+LF KM+I
                                                                                        Sbjct: 300 RKLASAFMNGYLRFDNQVIRNIAYELFRKMRI 331
Query= sid | 110162 | lan | 1820RF007 Phage 182 ORF | 7795-8775 | 1
         (326 letters)
>gi|1429239|emb|CAA67658| (X99260) upper collar protein
```

{Bacteriophage B103}

```
Length = 308
```

```
Score = 271 bits (685), Expect = 6e-72
Identities = 131/275 (47%), Positives = 187/275 (67%), Gaps = 5/275 (1%)
```

- Query: 36 YYEHYRRQLTLLTFQLFEWENLPKSIDPRYLEIALHTNGYLGFFKDPTLGFMVCAGAEDG 95 +Y HY + L L +QLFEWE LP S+DP YLE ++H GY+GF+KDP +G++ C GA G
- Sbjct: 22 WYYHYYQYLCSLAYQLFEWERLPPSVDPSYLEKSIHQFGYVGFYKDPRIGYIACQGALSG 81
- Query: 96 QIDHYHNPIFFTANEAMYHKRYPVLRYDDDDDKSKCIMLYNNDLKVPTLPSLHRFALDMA 155
- +DHY+ P F A+ Y + + Y D +K+ + +YNNDLK TLP+L FA D+A
  Sbjct: 82 TVDHYNLPDRFHASSVGYQNTFKLYNYSDMKEKNMGVAIYNNDLKCSTLPALEMFAQDLA 141
- Query: 156 DINQISRVNRRAQKTPVIIQTDEKKYFSLLQAYNQIDENNQAVFVDKDMEFDESFNVWQT 215 ++ +I VN+ AQKTPV+I ++ SL YNQ + N +FV + ++ D + V++T Sbjct: 142 ELKEIIAVNQNAQKTPVLIAANDNNQLSLKNIYNQYEGNAPVIFVHESLDLD-NLKVFKT 200
- Query: 216 NAPYVVDKLRSELNEVWNEVLTFLGINNANVDKTARVQTSEVLSNNEQIESSGNILLKSR 275 +APYVVDKL ++ N VWNEV+T+LGI NAN++K R+ TSEV SN+EQIESSGNI LK+R
- Sbjct: 201 DAPYVVDKLNAQKNAVWNEVMTYLGIKNANLEKKERMVTSEVDSNDEQIESSGNIYLKAR 260
- Query: 276 KEFCDRVNRVFGDELDGKIDVKFRTDAVRQLQLAA 310 +B C++++ ++G L VKFR D V Q++L A
- Sbjct: 261 QEACNKISELYGLNL----KVKFRYDIVEQMRLNA 291
- Query= sid|110163|lan|1820RF008 Phage 182 ORF|14105-14983|2 (292 letters)
- >gi|4210750|emb|CAA10710| (AJ132604) LysL protein [Lactococcus lactisl Length = 235
- Score = 139 bits (347), Expect = 2e-32 Identities = 85/210 (40%), Positives = 114/210 (53%), Gaps = 14/210 (6%)
- Query: 2 MNGIDISSYQTGIDLSKVPCDFVNIKATGGTGYVNPDCDRAFQQALSLGKKIGVYHFAHE 61 MNGIDISSYQ ++ VP DFV IKAT GT Y+NP + Q + K +G YHFA Sbjct: 1 MNGIDISSYQAELNAGIVPSDFVIIKATEGTNYINPTWEEQAGQVIQTNKLLGFYHFAS- 59
- Query: 62 RGLEGTPQQEAQFFLDNIKGYIGKAVLILDFEGS--NQKDVNWAKAFLDYVYNKTGVKAW 119
  G P EA FF+ +K YIGKAVL+LDFE N A+ FL+ V KTG+
  Sbjct: 60 ---VGNPIAEADFFISVVKNYIGKAVLVLDFEAGAINAWGNVGARQFLNRVKEKTGINPM 116
- Query: 120 FYTYTANLNTTDFSSIAKGDYGLWVAEYGSNQPQGYSQPAPPKTNN----FPIVACFQF 174
- Y + ++S+I+ + LWVA+Y S P GY + P T+ + A Q+
  Sbjct: 117 IYMSSDVTRQFNWSTISSTN-PLWVAQYASMNPTGYQ--SEPWTDGKGYGAWSSAAIHQY 173
- Query: 175 TSKGRLPGYNGNLDLNVFYGDGNTWDLYVG 204 +S G L ++GNLD+N+ Y + N W
- Sbjct: 174 SSAGSLSNWSGNLDINLAYINANQWKSLAG 203
- Query= sid|110164|lan|182ORF009 Phage 182 ORF|8765-9601|2 (278 letters)
- >gi|1429240|emb|CAA67659| (X99260) lower collar protein [Bacteriophage B103] Length = 293
- Score = 180 bits (451), Expect = 1e-44 Identities = 115/296 (38%), Positives = 161/296 (53%), Gaps = 33/296 (11%)
- LKRYIESFTYYOPELSRKERIEVGRKQLFDFDYPFYDETKRAEFETKFINHFYLREIGSE 62 L YIE ++ Y+ LS E+IE GR +LFDF YP +DE+ R FET FI +FY+REIG E
- LSTYIEMWSQYETGLSMAEKIEKGRPKLFDFQYPIFDESYRKVFETHFIRNFYMREIGFE 67
- Query: 63 TMGSPKFNLDEYLNLNMPYWNKMPLSNLEEF-PIFDDMDYTIDEKQKLLNEIDTNIKANR 121 T G FKFNL+ +L +NMPY+NK+F S L ++ P+ + T K+
- Sbjct: 68 TEGLFKFNLETWLIINMPYFNKLPESELIKYDPLENTRLNTTGNKKN-----DTERNDNR 122
- Query: 122 D-----ESKNQTKQVDQTDNRNKNTRDTGTT-----DSFSRNTYTDTPQKDLRIASNG 169 + K+ TK D+T+ + D TT D+F+R +D P L + +N

```
Sbjct: 123 DTTGSMKADGKSNTKTSDKTNATGSSKEDGKTTGSVTDDNFNRKIDSDQPDSRLNLTTN- 181
Query: 170 DGTGVINYATNITEDLSKETTSSTGVETNNDKTNQNTRSNAS-----EKETKNTD 219
DG G + YA+ I E+ + ++TG TNN ++ + S S T N
Sbjct: 182 DGQGTLEYASAIEENNTNNKRNTTG--TNNVTSSAESESTGSGTSDTVTTDNANTTTNDK 239
Query: 220 INKDQNQTKDTITRYKGKKGNTDYADLLEKYRRSVLRIEKMIFREMNKEGLFLLVY 275
+N N +D I GK G YA L++ YR ++LRIEK IF EM + LF+LVY
Sbjct: 240 LNSQINNVEDYIESKIGKSGTQSYASLVQDYRAALLRIEKRIFDEMQE--LFMLVY 293
Query= sid|110165|1an|182ORF010 Phage 182 ORF|1310-2155|2
         (281 letters)
>gi|135604|sp|P06812|TERM_BPNF DNA TERMINAL PROTEIN
           >gi|75815|pir||ERBPNP terminal protein - phage NF
>gi|579177|emb|CAA68440| (Y00363) gene E product (AA
           1-267) [Bacteriophage NF]
          Length = 266
 Score = 74.9 bits (181), Expect = 6e-13
 Identities = 73/275 (26%), Positives = 129/275 (46%), Gaps = 37/275 (13%)
         VRISKNDRAKLEKIYGKSNKARKKYNRLRQK-GVE---ERQLPTVPTSKKRLIDYVKSTN 58
           +RI+ ND+A K+ K+ KA K +R ++K G++ E +LP + + +
          IRITNNDKALYAKLV-KNTKA--KISRTKKKYGIDLSNEIELPPLESFQ----- 52
Sbict: 7
Query: 59 MSRSDFNKMLDELVDFAQPYNENYIFEINKRNVAISRAQIKEAQIKTEQAQKAKEEHYKE 118
            +R +FNK + F N+NY F NK + S+A+I E T++AQ+ +E +E
Sbjct: 53 -TREEFNKWKQKQESFTNRANQNYQFVKNKYGIVASKAKINEIAKNTKEAQRIVDEQREE 111
I++P+ +T G P DFN D S +++ E
                   K +
Sbjct: 112 IEDKPFISGGKQQGTVGQRMQILSPSQVT--GISRP----SDFNFDDVRSYARLRTLEEG 165
Query: 171 IG-KQDEQYFDERDQLYYDNFRQAMFTIFNSD--ADDIVRLLDSMGLDLFMKTYVSNFLD 227
           + K Y+D R + NF + + FNSD +D++V L + D F + Y+ F +
Sbjct: 166 MAEKASPDYYDRRMTQMHQNFIEIVEKSFNSDWLSDELVERLKKIPPDDFFELYLM-FDE 224
Query: 228 MNLDYIYDEAEVQQKKEQVYSKIAKVIESETGGEV 262
           ++ +Y E E + E + +KI ++
                                         G+V
Sbjct: 225 ISFEYFDSEGEDVEASEAMLNKIHSYLDRYERGDV 259
Query= sid|110166|lan|1820RF011 Phage 182 ORF|9607-10158|1
         (183 letters)
>gi|1429241|emb|CAA67660| (X99260) pre-neck appendage protein
          [Bacteriophage B103]
          Length = 860
 Score = 50.8 bits (119), Expect = 6e-06
 Identities = 29/105 (27%), Positives = 56/105 (52%), Gaps = 6/105 (5%)
          KRFDGLPAVFKERFSKYPHTEYRYELLLDEEVSALIAYLNEVGALVNDMSGYLNYFIEHF 67
           +RF+ L + + + +Y T + + L E+++ +I YLN++G L ND+
          RRFEKLGEMMVQVYERYLPTAFDESMTLLEKMNKIIEYLNQIGRLTNDVVEEWNKVMEWI 66
Query: 68 V-EKLEEITNDTLKKWLSDGTLENLINDTVFANYIKEIKRLQILV 111
                                        I E+K+
            + LE+ +TL+KW +G +L+
Sbjct: 67 LNDGLEDYVKETLEKWYEEGKFADLV----IQVIDELKQFGVSV 106
                                                                              - ______
Query= sid | 110169 | lan | 1820RF014 Phage 182 ORF | 13716-14108 | 3
         (130 letters)
>gi|137936|sp|P11188|VG14_BPPH2 LYSIS PROTEIN (LATE PROTEIN GP14)
          >gi|75860|pir||WMBP29 gene 14 protein - phage phi-29
```

>gi|15678|emb|CAA28631| (X04962) gene 14 product (AA

```
1-393) [Bacteriophage phi-29] >gi|225369|prf||1301270J
           gene 14 [Bacteriophage phi-29]
           Length = 131
 Score = 96.7 bits (237), Expect = 6e-20
 Identities = 53/131 (40%), Positives = 81/131 (61%), Gaps = 3/131 (2%)
Query: 1 MIEYITQWL-ADDNHLVYGLIIWLMVAMIIDFVLGFTIAKFNKEIDFSSFKAKAGIIVKV 59
           MI ++ +L D+ L+Y L +LMV M++D VLG AK N I FSSFK K G+++KV
           MIAWMOHFLETDETKLIYWLT-FLMVCMVVDTVLGVLFAKLNPNIKFSSFKIKTGVLIKV 61
Sbict: 3
Query: 60 AEMVLVVYFIPVAVKFGAVGITMYITMLVGLILSEIYSILGHISDIDDDNNWTDYVKKFL 119
           +EM+L + IP AV F A G+ + T+ L +SEIYSI GH+ +DD +++ + ++ F
Sbjct: 62 SEMILALLAIPFAVPFPA-GLPLLYTVYTALCVSEIYSIFGHLRLVDDKSDFLEILENFF 120
Query: 120 DGTLNRKDDIK 130
Sbjct: 121 KRTSGKNKEEK 131
Query= sid|110170|lan|1820RF015 Phage 182 ORF|854-1225|2
         (123 letters)
>gi|15670|emb|CAA24483| (V01155) reading frame 10 (may be gene 4)
           [Bacteriophage phi-29]
           Length = 124
 Score = 69.9 bits (168), Expect = 6e-12
 Identities = 39/119 (32%), Positives = 64/119 (53%), Gaps = 3/119 (2%)
           IVKSTFDTQTPEGMLQVFNATNGASIPLRNAI-GEVLELKDILVYSDEVSGFGGAEPSQA 61
           IVK+TFDT+T EG +++FNA G +N G ++E I Y
           IVKATFDTETLEGQIKIFNAQTGGGQSFKNLPDGTIIEANAIAQYKQVSDTYGDAK--EE 63
Sbict: 6
Query: 62 ELVAFFTEDGKTYAGVSAVATKSAKNLIDMMTANPDIKPKISFVEGKSNGGQKFVNLQV 120
+ F DG Y+ +S ++A +LID++T + K+ V+G S+ G F +LQ+
Sbjct: 64 TVTTIFAADGSLYSAISKTVAEAASDLIDLVTRHKLETFKVKVVQGTSSKGNVFFSLQL 122
Query= sid | 110174 | lan | 1820RF019 Phage 182 ORF | 4323-4613 | 3
         (96 letters)
>gi|1429235|emb|CAA67654| (X99260) head morphogenesis protein
          [Bacteriophage B103]
 Score = 60.9 bits (145), Expect = 1e-09
 Identities = 34/96 (35%), Positives = 53/96 (54%), Gaps = 5/96 (5%)
Query: 1 MEIKEHESILNGILESVTDGEARSKIVEHLEALREDYGATTEALTSANSTLEKLKKDNEA 60
MB HE ILN + + + R+++ L+ LR DYG+ + S EKL+ +N
Sbjct: 3 MERDSHEEILNKLNDPELEHSERTEL---LQQLRADYGSVLSEFSELTSATEKLRAENSD 59
Query: 61 LVISNSKLFRERAIVEPAEN--NEPETDQNITLDDL 94
          L++SNSKLFR+ I + E + E + IT++DL
Sbjct: 60 LIVSNSKLFRQVGITKEKEEBIKQEELSETITIEDL 95
Query= sid|110180|lan|182ORF025 Phage 182 ORF|548-814|2
         (88 letters)
                                                                                 >gi|138099|sp|P06955|VG6_BPPZA EARLY PROTEIN GP6
          >gi|75841|pir||ERBP6Z gene 6 protein - phage PZA
>gi|216047 (M11813) gene 6 product [Bacteriophage PZA]
          >gi|224746|prf||1112171K ORF 6 [Bacteriophage PZA]
          Length = 96
 Score = 55.0 bits (130), Expect = 8e-08
 Identities = 28/79 (35%), Positives = 45/79 (56%)
```

Query: 4 KLMQRNVTSTKVEFSEVIVQDGAPTIVPCEPVVLTGKLSEEKALSAIKRKNPDKNVVVTN 63
K+MQR +T T V +++++ DG + G LS E+A +KRK + V V +
Sbjgt: 3 KMMQREITKTTVNVAKMVMVDGEVQVEQLPSETFVGNLSMEQAQWRMKRKYKGEPVQVVS 62

Query: 64 VSHETALYTMPVDKFIELA 82

V T +Y +PV+KF+E+A
Sbjct: 63 VEPNTEVYELPVEKFLEVA 81

## Table 26

## Secondary structure prediction for ORF 1820RF008

1	MMNGIDISSY	QTGIDLSKVP	CDFVNIKATG	GTGYVNPDCD	RAFQQALSLG	KKIGVYHFAH
	CCCCCCCCC	CCCCCCCCC	CCEEEEECC	CCCCCCCCC	нинининис	CCCCEEEEE
61	ERGLEGTPQQ	EAQFFLDNIK	GYIGKAVLIL	DFEGSNQKDV	NWAKAFLDYV	YNKTGVKAWF
	CCCCCCCHH	нинининис	CCCCEEEEE	ССССССННН	нинининин	HCCCCCEEEE
121	YTYTANLNTT	DFSSIAKGDY	GLWVAEYGSN	QPQGYSQPAP	PKTNNFPIVA	CFQFTSKGRL
	EEECCCCCCC	CCCEECCCCC	CEEEEECCCC	CCCCCCCCC	CCCCCCEEE	EEEECCCCCC
181	PGYNGNLDLN	VFYGDGNTWD	LYVGKKQDQI	VPPENKIFDA	TSDEFIFTLT	TGSTSVFYFD
	CCCCCCCEE	EEECCCCCCE	EEECCCCCC	CCCCCCCCC	CCCEEEEEC	CCCCEEECC
241	GETIFELSDP	TQLDHIRGTY	NHVHGKEIPS	MVWTPEQFDI	YLKMYEKKPV	YK
	CCEEEECCCC	CCHHHHCCEE	CCCCCEECC	ССССССИНН	ННННССССЕ	EC

## Secondary structure prediction for ORF 1820RF014

- 1 MIEYITQWLA DDNHLVYGLI IWLMVAMIID FVLGFTIAKF NKEIDFSSFK AKAGIIVKVA ССССЕЕСССС ССССИННИН НИНИНИНИ ИНИНИНИНИ СССССИНИНИ НИНСЕВЕВЕ
- 61 EMVLVVYFIP VAVKFGAVGI TMYITMLVGL ILSEIYSILG HISDIDDDNN WTDYVKKFLD
- 121 GTLNRKDDIK

CCCCCCEEC

#### Table 27

#### Enterococcus accession numbers 242/242

gi|2895751|gb|AF044978.1|AF044978 [2895751] gi|4803755|dbj|AB026843.1|AB026843 [4803755] gi|4769001|gb|AF140549.1|AF140549 [4769001] gi|4760901|gb|AF099088.1|AF099088 [4760901] gi|4704705|gb|AF121254.1|AF121254 [4704705] gi|3342117|gb|AF076604.1|AF076604 [3342117] gi|4688824|emb|AJ132470.1|ESP132470 [4688824] gi|4732085|gb|AF125553.1|AF125553 [4732085] gi|4732082|gb|AF125552.1|AF125552 [4732082] gi|4732079|gb|AF125551.1|AF125551 [4732079] gi|4732076|gb|AF125550.1|AF125550 [4732076] gi|4732073|gb|AF125548.1|AF125548 [4732073] gi|4732070|gb|AF125547.1|AF125547 [4732070] gi|4732067|gb|AF125546.1|AF125546 [4732067] gi|4732064|gb|AF125545.1|AF125545 [4732064] gi|4732061|gb|AF125544.1|AF125544 [4732061] gi|4704653|gb|AF114715.1|AF114715 [4704653] gi|4704564|gb|AF102550.1|AF102550 [4704564] gi|4688827|emb|AJ238249.1|EFA238249 [4688827] gi|4680606|gb|AF125198.1|AF125198 [4680606] gi|4633279|gb|AF117609.1|AF117609 [4633279] gi|4633124|gb|AF110130.1|AF110130 [4633124] gi|4590399|gb|AF124258.1|AF124258 [4590399] gi|4590336|gb|AF108380.1|AF108380 [4590336] gi|4590335|gb|AF108379.1|AF108379 [4590335] gi|4019167|gb|U21300.1|CXU21300 [4019167] gi|4545122|gb|AF077816.1|AF077816 [4545122] gi|4433610|gb|AF106614.1|AF106614 [4433610] gi|4468838|emb|AJ132039.1|EFA132039 [4468838] gi|4468121|emb|AJ132958.1|BPH132958 [4468121] gi|4456104|emb|Y17302.1|EHI17302 [4456104] gi|4433611|gb|AF106615.1|AF106615 [4433611]

gi|4433607|gb|AF106611.1|AF106611 [4433607]

gi|4098267|gb|U76614.1|BLU76614 [4098267] gi|47019|emb|Y00116.1|SFAMB1 [47019] gi|4158179|emb|AL035206.1|SC9B5 [4158179] gi|4165458|emb|X79343.1|EF16SSPA [4165458] gi|4165457|emb|X79342.1|EFTRNALA [4165457] gi|4165456|emb|X79341.1|EF23SRNA [4165456] gi|4150978|emb|Y14027.1|EFY14027 [4150978] gi|4127803|emb|AJ223161.1|EFAJ3161 [4127803] gi|2956685|emb|Y16413.1|EFENTIJO [2956685] gi|2665346|emb|Y13922.1|EHY13922 [2665346] gi|4324675|gb|AF109375.1|AF109375 [4324675] gi|4234627|gb|AF061013.1|AF061013 [4234627] gi|4234626|gb|AF061012.1|AF061012 [4234626] gi|4234625|gb|AF061011.1|AF061011 [4234625] gi|4234624|gb|AF061010.1|AF061010 [4234624] gi|4234623|gb|AF061009.1|AF061009 [4234623] gi|4234622|gb|AF061008.1|AF061008 [4234622] gi|4234621|gb|AF061007.1|AF061007 [4234621] gi|4234620|gb|AF061006.1|AF061006 [4234620] gi|4234619|gb|AF061005.1|AF061005 [4234619] gi|4234618|gb|AF061004.1|AF061004 [4234618] gi|4234617|gb|AF061003.1|AF061003 [4234617] gi|4234616|gb|AF061002.1|AF061002 [4234616] gi|4234615|gb|AF061001.1|AF061001 [4234615] gi|4234614|gb|AF061000.1|AF061000 [4234614] gi|3138990|gb|AF060241.1|AF060241 [3138990] gi|3138986|gb|AF060240.1|AF060240 [3138986] gi|4204535|gb|AF094803.1|AF094803 [4204535] gi|4204534|gb|AF094802.1|AF094802 [4204534] gi|4204533|gb|AF094801.1|AF094801 [4204533] gi|4204532|gb|AF094800.1|AF094800 [4204532] gi|4204531|gb|AF094799.1|AF094799-{4204531}gi|4204530|gb|AF094798.1|AF094798 [4204530] gi|4204529|gb|AF094797.1|AF094797 [4204529] gi|4204528|gb|AF094796.1|AF094796 [4204528] gi|4204527|gb|AF094795.1|AF094795 [4204527]

gi|4204526|gb|AF094794.1|AF094794 [4204526] gi|4204525|gb|AF094793.1|AF094793 [4204525] gi|4204524|gb|AF094792.1|AF094792 [4204524] gi|4204523|gb|AF094791.1|AF094791 [4204523] gi|4204522|gb|AF094790.1|AF094790 [4204522] gi|4204521|gb|AF094789.1|AF094789 [4204521] gi|4204520|gb|AF094788.1|AF094788 [4204520] gi|4204519|gb|AF094787.1|AF094787 [4204519] gi|4204518|gb|AF094786.1|AF094786 [4204518] gi|4204517|gb|AF094785.1|AF094785 [4204517] gi|4204516|gb|AF094784.1|AF094784 [4204516] gi|4204515|gb|AF094783.1|AF094783 [4204515] gi|4204514|gb|AF094782.1|AF094782 [4204514] gi|4204513|gb|AF094781.1|AF094781 [4204513] gi|4204512|gb|AF094780.1|AF094780 [4204512] gi|3873186|gb|AF034779.1|AF034779 [3873186] gi|4151367|gb|AF093508.1|AF093508 [4151367] gi|2828136|gb|AF039903.1|AF039903 [2828136] gi|2828135|gb|AF039902.1|AF039902 [2828135] gi|2828134|gb|AF039901.1|AF039901 [2828134] gi|2828133|gb|AF039900.1|AF039900 [2828133] gi|2828132|gb|AF039899.1|AF039899 [2828132] gi|2828131|gb|AF039898.1|AF039898 [2828131] gi|4103866|gb|AF028812.1|AF028812 [4103866] gi|4103864|gb|AF028811.1|AF028811 [4103864] gi|2605925|gb|AF029727.1|AF029727 [2605925] gi|1402750|gb|U60038.1|EFU60038 [1402750] gi|1835780|gb|U86375.1|EFU86375 [1835780] gi|3831555|gb|AF047608.1|AF047608 [3831555] gi|3790617|gb|AF097414.1|AF097414 [3790617] gi|3767587|dbj|AB005036.1|AB005036 [3767587] gi|3757810|gb|AF042288.1|AF042288 [3757810] gi|3747039|gb|AF093509.1|AF093509 [3747039] gi|3660559|dbj|AB017811.1|AB017811 [3660559] gi|1147743|gb|U42211.1|EHU42211 [1147743] gi|3676412|gb|AF051917.1|AF051917 [3676412] gi|3676164|emb|AJ011113.1|EFA011113 [3676164] gi|2612869|gb|AF005726.1|AF005726 [2612869] gi|2353762|gb|AF016233.1|AF016233 [2353762]

gi|2149899|gb|U94707.1|EFU94707 [2149899] gi|2149149|gb|U82366.1|LSU82366 [2149149] gi|1469463|gb|U49512.1|EFU49512 [1469463] gi|1244503|gb|U35366.1|EFU35366 [1244503] gi|833854|gb|U26268.1|EFU26268 [833854] gi|841200|gb|U18931.1|CPU18931 [841200] gi|460079|gb|U00457.1|U00457 [460079] gi|460077|gb|U00456.1|U00456 [460077] gi|535661|gb|L34675.1|INSTRANSPO [535661] gi|3023041|gb|AF007787.1|AF007787 [3023041] gi|431124|gb|L15633.1|TRN916ENT [431124] gi|388106|gb|L23802.1|ENEEBSA [388106] gi|3608387|gb|AF071085.1|AF071085 [3608387] gi|3551851|gb|AF076027.1|AF076027 [3551851] gi|3551773|gb|U94770.1|SPU94770 [3551773] gi|3551743|gb|U57498.1|ECU57498 [3551743] gi|3243178|gb|AF063010.1|AF063010 [3243178] gi|3136316|gb|AF063900.1|AF063900 [3136316] gi|3540256|gb|AF052459.1|AF052459 [3540256] gi|755215|gb|U17696.1|LLU17696 [755215] gi|3421437|gb|AF082295.1|AF082295 [3421437] gi|3421436|gb|AF082294.1|AF082294 [3421436] gi|3421435|gb|AF082293.1|AF082293 [3421435] gi|3421434|gb|AF082292.1|AF082292 [3421434] gi|3341430|emb|Y17797.1|EFY17797 [3341430] gi|3319647|emb|X69092.1|EHPBP3RA [3319647] gi|3292886|emb|AJ007584.1|EFA7584 [3292886] gi|3261536|emb|AL021958.1|MTV041 [3261536] gi|3250708|emb|Z95150.1|MTCY164 [3250708] gi|3249688|gb|AF070678.1|AF070678 [3249688] gi|3249687|gb|AF070677.1|AF070677 [3249687] gi|3249686|gb|AF070676.1|AF070676 [3249686] gi|3219158|dbj|AB015233.1|AB015233 [3219158] gi|2765275|emb|Y12924.1|SPY12924 [2765275] gi|3183687|emb|Y11621.1|EA16SRRN [3183687] gi|2765274|emb|Y12923.1|EFY12923 [2765274] = gi|2765273|emb|Y12922.1|ESY12922 [2765273] gi|2765272|emb|Y12921.1|ESY12921 [2765272] gi|2765271|emb|Y12920.1|EDY12920 [2765271] gi|2765270|emb|Y12919.1|ESY12919 [2765270]

gi|2058762|gb|B07882.1|B07882 [2058762] gi|2765269|emb|Y12918.1|ECY12918 [2765269] gi|2058761|gb|B07881.1|B07881 [2058761] gi|2765268|emb|Y12917.1|ECY12917 [2765268] gi|2058760|gb|B07880.1|B07880 [2058760] gi|2765267|emb|Y12916.1|EPY12916 [2765267] gi|2765266|emb|Y12915.1|ESY12915 [2765266] gi|2058759|gb|B07879.1|B07879 [2058759] gi|2765265|emb|Y12914.1|ERY12914 [2765265] gi|2058758|gb|B07878.1|B07878 [2058758] gi|2058757|gb|B07877.1|B07877 [2058757] gi|2765264|emb|Y12913.1|EMY12913 [2765264] gi|2765263|emb|Y12912.1|EHY12912 [2765263] gi|2058756|gb|B07876.1|B07876 [2058756] gi|2058755|gb|B07875.1|B07875 [2058755] gi|2765262|emb|Y12911.1|EMY12911 [2765262] gi|2765261|emb|Y12910.1|EGY12910 [2765261] gi|2058754|gb|B07874.1|B07874 [2058754] gi|2058753|gb|B07863.1|B07863 [2058753] gi|2765260|emb|Y12909.1|EDY12909 [2765260] gi|2765259|emb|Y12908.1|ECY12908 [2765259] gi|2058752|gb|B07862.1|B07862 [2058752] gi|2765258|emb|Y12907.1|EAY12907 [2765258] gi|2058751|gb|B07861.1|B07861 [2058751] gi|2058750|gb|B07860.1|B07860 [2058750] gi|2765257|emb|Y12906.1|EFY12906 [2765257] gi|2058749|gb|B07859.1|B07859 [2058749] gi|2765256|emb|Y12905.1|EFY12905 [2765256] gi|2894541|emb|AJ223332.1|EFAJ3332 [2894541] gi|2058748|gb|B07858.1|B07858 [2058748] gi|2058747|gb|B07857.1|B07857 [2058747] gi|2894539|emb|AJ223331.1|EFAJ3331 [2894539] gi|3108058|gb|AF060881.1|AF060881 [3108058] gi|2058746|gb|B07856.1|B07856 [2058746] gi|2058745|gb|B07855.1|B07855 [2058745] gi|3087776|emb|AJ223633.1|EFAJ3633 [3087776] gi|3080754|gb|AF016483.1|AF016483 [3080754] gi|2058744|gb|B07854.1|B07854 [2058744] gi|2058743|gb|B07853.1|B07853 [2058743] gi|2197119|gb|AF003921.1|AF003921 [2197119] gi|2058742|gb|B07852.1|B07852 [2058742] gi|2982722|dbj|AB012213.1|AB012213 [2982722] gi|2058741|gb|B07851.1|B07851 [2058741] gi|2982721|dbj|AB012212.1|AB012212 [2982721] gi|2058740|gb|B07850.1|B07850 [2058740] gi|2058780|gb|B07890.1|B07890 [2058780] gi|2947527|gb|T25933.1|T25933 [2947527] gi|2058779|gb|B07889.1|B07889 [2058779] gi|2924302|emb|X81655.1|EHERMAM [2924302] gi|2058778|gb|B07888.1|B07888 [2058778] gi|2664256|emb|Y12234.1|EFAS48C [2664256] gi|2058777|gb|B07887.1|B07887 [2058777] gi[2879906|dbj[D85752.1|D85752 [2879906] gi|2058776|gb|B07886.1|B07886 [2058776] gi|2746216|gb|AF028836.1|AF028836 [2746216] gij2058775|gb|B07885.1|B07885 [2058775] gi|2745825|gb|AF039139.1|AF039139 [2745825] gi|2058774|gb|B07884.1|B07884 [2058774] gi|2696019|dbj|AB007844.1|AB007844 [2696019] gi|2058773|gb|B07873.1|B07873 [2058773] gi|48999|emb|X62280.1|EHPBP5G [48999] gi|2058772|gb|B07872.1|B07872 [2058772] gi|2654477|gb|U89914.1|BFU89914 [2654477] gi|2058771|gb|B07871.1|B07871 [2058771] gi|43347|emb|X68646.1|EHPSRAA [43347] gi|2058770|gb|B07870.1|B07870 [2058770] gi|2613034|gb|AH005624.1|SEG\_EDDH4RR gi|2058769|gb|B07869.1|B07869 [2058769] [2613034] gi|2058768|gb|B07868.1|B07868 [2058768] gi|2613033|gb|AF029775.1|EDDH4RR2 [2613033] gi|2058767|gb|B07867.1|B07867 [2058767] gi|2613032|gb|AF029774.1|EDDH4RRI [2613032] gi|2058766|gb|B07866.1|B07866 [2058766] gi|2613031|gb|AH005623.1|SEG\_EDDHIRR gi|2058765|gb|B07865.1|B07865 [2058765] [2613031] gi|2058764|gb|B07864.1|B07864 [2058764] gi|2613030|gb|AF029773.1|EDDHIRR2 [2613030] gi|2058763|gb|B07883.1|B07883 [2058763]

gi|2613029|gb|AF029772.1|EDDHIRR1 [2613029] gi|2613028|gb|AH005622.1|SEG\_EDH19RR [2613028] gi|2613027|gb|AF029771.1|EDH19RR2 [2613027] gi|2613026|gb|AF029770.1|EDH19RR1 [2613026] gi|2613025|gb|AH005621.1|SEG\_EDISRR [2613025] gi|2613024|gb|AF029769.1|EDISRR2 [2613024] gi|2613023|gb|AF029768.1|EDISRR1 [2613023] gi|1881226|dbj|AB001488.1|AB001488 [1881226] gi|2547160|gb|AF023104.1|AF023104 [2547160] gi|2547159|gb|AF023103.1|AF023103 [2547159] gi|2547158|gb|AF023102.1|AF023102 [2547158] gi|2547157|gb|AF023101.1|AF023101 [2547157] gi|2415383|gb|AF015775.1|AF015775 [2415383] gi|2388636|gb|U94356.1|EFU94356 [2388636] gi|2388634|gb|U94355.1|ECU94355 [2388634] gi|2340825|dbj|D26045.1|D26045 [2340825] gi|2226147|emb|Y14080.1|BSY14080 [2226147] gi|2327026|gb|U87997.1|EFU87997 [2327026] gi|2318058|gb|AF012532.1|AF012532 [2318058] gi|1848175|emb|X87189.1|EM23S5SSP [1848175] gi|1848174|emb|X87187.1|EM16S23SS [1848174] gi|1848173|emb|X87188.1|EM16S23SP [1848173] gi|1848172|emb|X87185.1|EH23S5SSP [1848172] gi|1848171|emb|X87184.1|EH16S23SS [1848171] gi|1848170|emb|X87181.1|EF23S5SSP [1848170] gi|1848169|emb|X87183.1|EF23S5SPA [1848169] gi|1848168|emb|X87191.1|EF23S5SAC [1848168] gi|1848167|emb|X87180.1|EF16S23SS [1848167] gi|1848166|emb|X87182.1|EF16S23SP [1848166] gi|1848165|emb|X87190.1|EF16S23SC [1848165] gi|1848164|emb|X87186.1|EF16S23SA [1848164] gi|1848156|emb|X87179.1|ED23S5SSP [1848156] gi|1848155|emb|X87178.1|ED16S23SS [1848155] gi|1848154|emb|X87177.1|ED16S23SA [1848154] gi|2274942|emb|AJ000346.1|EHNAPBC [2274942] gi|2274939|emb|AJ000042.1|EFGLS24B [2274939] gi|414575|gb|L12710.1|ENEAAC [414575] gi|2245603|gb|AF006008.1|AF006008 [2245603]

gi|2231992|gb|U94530.1|EFU94530 [2231992] gi|2231990|gb|U94529.1|EFU94529 [2231990] gi|2231988|gb|U94528.1|EFU94528 [2231988] gi|2231986|gb|U94527.1|EFU94527 [2231986] gi|2231984|gb|U94526.1|EFU94526 [2231984] gi|2231982|gb|U94525.1|ECU94525 [2231982] gi|2231980|gb|U94524.1|ECU94524 [2231980] gi|2231978|gb|U94523.1|ECU94523 [2231978] gi|2231976|gb|U94522.1|ECU94522 [2231976] gi|2231974|gb|U94521.1|ECU94521 [2231974] gi|2196685|gb|U25090.1|EFU25090 [2196685] gi|2197120|gb|AF003922.1|AF003922 [2197120] gi|2196683|gb|U25095.1|EFU25095 [2196683] gi|2196681|gb|U25094.1|EFU25094 [2196681] gi|2196679|gb|U25093.1|EFU25093 [2196679] gi|2196677|gb|U25092.1|EFU25092 [2196677] gi|2196675|gb|U25091.1|EFU25091 [2196675] gi|2196673|gb|U24682.1|EFU24682 [2196673] gi|532533|gb|U09422.1|EFU09422 [532533] gi|487271|dbi|D17462.1|ENENTP [487271] gi|468459|dbi|D28859.1|ENEPPD1 [468459] gi|440135|dbj|D16334.1|ENEATPK [440135] gi|391680|dbj|D13816.1|ENENAABS [391680] gi|1402524|dbj|D78257.1|D78257 [1402524] gi|709995|dbj|D30808.1|BACYCB20 [709995] gi|2109265|gb|U91527.1|EFU91527 [2109265] gi|1041112|dbj|D78016.1|ENEPPD1A [1041112] gi|1339880|dbj|D85392.1|ENERPA [1339880] gi|1339878|dbj|D85393.1|ENEGE1E [1339878] gi|662918|emb|Z46807.1|EHCOPAYZ [662918] gi|769796|emb|X86176.1|EFRPODDNE [769796] gi|1854638|gb|U51479.1|EGU51479 [1854638] gi|1857221|gb|U72706.1|EFU72706 [1857221] gi|1857219|gb|U72704.1|EFU72704 [1857219] gi|1857217|gb|U72705.1|ECU72705 [1857217] gi|1272655|emb|X96978.1|EFPPD1GNS [1272655] gi|1272652|emb|X96976.1|EFPLSEPIG [1272652] gi|1279406|emb|X96977.1|EFPAD1ORF [1279406] gi|1070149|emb|X93211.1|EFTNFO1 [1070149]

gi|1065723|emb|X92947.1|EFTETMGN [1065723] gi|1019639|gb|L38972.1|PH4COINJN [1019639] gi|1151151|gb|U43087.1|EFU43087 [1151151] gi|1098507|gb|U17283.1|BMU17283 [1098507] gi|1498072|gb|U64887.1|EFU64887 [1498072] gi|1498071|gb|U64886.1|EFU64886 [1498071] gi|1469783|gb|U58049.1|EHU58049 [1469783] gi|1763666|gb|U81452.1|EFU81452 [1763666] gi|624694|gb|L38973.1|PH4SEQ [624694] gi|1730458|emb|Z83305.1|EFVANRES [1730458] gi|1419498|emb|X84796.1|ECPFW4 [1419498] gi|1419497|emb|X84795.1|ECPFW3 [1419497] gi[1419496]emb[X84794.1]ECPFW1 [1419496] gi|254400|gb|S43266.1|S43266 [254400] gi|239025|gb|S66277.1|S66277 [239025] gi|1054931|gb|U38590.1|EFU38590 [1054931] gi|1244573|gb|U39788.1|EHU39788 [1244573] gi|1244571|gb|U39789.1|EGU39789 [1244571] gi|1244569|gb|U39790.1|EFU39790 [1244569] gi|1255020|gb|U39777.1|ESU39777 [1255020] gi|1255018|gb|U39775.1|EPU39775 [1255018] gi|1255016|gb|U39778.1|EDU39778 [1255016] gi|1255014|gb|U39776.1|ECU39776 [1255014] gi|1255012|gb|U39774.1|EAU39774 [1255012] gi|1619922|gb|U69267.1|IVU69267 [1619922] gi|790436|emb|X84861.1|EFEFMPBP5 [790436] gi|790434|emb|X84858.1|EFD63RPSR [790434] gi|790432|emb|X84862.1|EF721PBP5 [790432] gi|790430|emb|X84860.1|EF63RPBP5 [790430] gi|790428|emb|X84859.1|EF366PBP5 [790428] gi|1572800|gb|U70854.1|CELF38A5 [1572800] gi|1041816|gb|U17153.1|EFU17153 [1041816] gi|1086523|gb|U39859.1|EFU39859 [1086523] gi|403564|gb|U01917.1|EFU01917 [403564] gi|1515474|gb|U66286.1|EFU66286 [1515474] gi|1513068|gb|U15554.1|LMU15554 [1513068] gi|1296520|emb|X94181.1|EFENTAORF [1296520] gi|1488069|gb|U63997.1|EFU63997 [1488069] gi|1209525|gb|U35369.1|EFU35369 [1209525]

gi|1469341|gb|U30931.1|ESU30931 [1469341] gi|488331|gb|M77276.1|SYNGIP2122 [488331] gi|1046177|gb|U39733.1| [1046177] gi|1236613|gb|U49939.1|CVU49939 [1236613] gi|47491|emb|X55766.1|SS16SR5G [47491] gi|47490|emb|X55767.1|SS16SR3G [47490] gi|47061|emb|X56353.1|SFTET916 [47061] gi|49022|emb|X62755.1|SFNPRG [49022] gi|47047|emb|X17214.1|SFPASA1 [47047] gi|47044|emb|X68847.1|SFNOXAA [47044] gi|47033|emb|V01547.1|SFKANR [47033] gi|47018|emb|X02027.1|SF5SRNA [47018] gi|511044|emb|X75752.1|MP16SRNA0 [511044] gi|511043|emb|X75751.1|MP16SR243 [511043] gi|886481|emb|X82819.1|ESPLPAM [886481] gi|517387|emb|X76177.1|ES16SRR [517387] gi|472916|emb|X76913.1|EHNTPOP [472916] gi|43351|emb|X55133.1|ES16SRRN [43351] gi|1143442|emb|X92687.1|EFPBP5G [1143442] gi|963032|emb|Z50854.1|EHARPQTOU [963032] gi|886479|emb|X84818.1|EHDNAPSR [886479] gi|551437|emb|X81654.1|EHIS1216 [551437] gi|467805|emb|X78425.1|EFPBP5 [467805] gi|296721|emb|X55961.1|EFPD78 [296721] gi|287946|emb|Z19137.1|EFPTSHGN [287946] gi|49042|emb|X63285.1|EHNAKA [49042] gi|49019|emb|X62658.1|EFSEA1 [49019] gi|43337|emb|Z12296.1|EFSPREG [43337] gi|43335|emb|X56895.1|EFPVANAG [43335] gi|43333|emb|X16421.1|EFPF54 [43333] gi|43331|emb|X62657.1|EFORF3 [43331] gi|1065721|emb|X92945.1|EFCAT501 [1065721] gi|806551|emb|Z49243.1|EF4110SOD [806551] gi|806549|emb|Z49244.1|EF4105SOD [806549] gi|505530|emb|X79542.1|EFAS48 [505530] gi|43323|emb|X62656.1|EFASP1-[43323] gi|40840|emb|X56422.1|EC16SRNAG [40840] gi|48189|emb|X04388.1|TN1545TR [48189] gi|928814|gb|L40841.1|ENETRANSPO [928814] gi|141856|gb|L01794.1|AD1REPABC [141856]

gi|149125|gb|M90647.1|IP8VANY [149125] gi|141862|gb|M87836.1|AD1TRAE1 [141862] gi|141860|gb|M84374.1|AD1TRAA [141860] gi|141853|gb|M62888.1|AD1PAD1 [141853] gi|1101637|dbj|D31674.1|EVM16RNA7 [1101637] gi|1101636|dbj|D31675.1|ENE16RNA8 [1101636] gi|497792|dbj|D31676.1|ENC16RNA9 [497792] gi|1022729|gb|U36195.1|EFU36195 [1022729] gi|488338|gb|M77279.1|SYNGIP3124 [488338] gi|488335|gb|M77278.1|SYNGIP2563 [488335] gi|488333|gb|M77277.1|SYNGIP2124 [488333] gi|488329|gb|M77275.1|SYNGIP2121 [488329] gi|388267|gb|L19532.1|AD1TRAC [388267] gi|493016|gb|U03756.1|EFU03756 [493016] gi|453536|gb|L28754.1|INSTRAN [453536] gi|153658|gb|M58002.1|STRHYDROLA [153658] gi|475427|gb|U00681.1|EFU00681 [475427] gi[818704|gb|U24692.1|EFU24692 [818704] gi|155036|gb|M97297.1|TRNVAN [155036] gi|150552|gb|M64978.1|PCFPRGAB [150552] gi|786274|gb|U22541.1|EHU22541 [786274] gi|786273|gb|U22540.1|EHU22540 [786273] gi|559858|gb|L37110.1|AD1CLYL [559858] gi|643614|gb|U16659.1|ECU16659 [643614] gi|643612|gb|U16658.1|ECU16658 [643612] gi|290641|gb|L13292.1|ENECOPPUMP [290641] gi|624701|gb|L29639.1|ENEVANCRF [624701] gi|624699|gb|L29638.1|ENEVANCR [624699] gi|624692|gb|L29641.1|ENEDDLA [624692] gi|624690|gb|L29640.1|ENEDDL [624690] gi|493094|gb|L32813.1|ENERRD [493094]

gi|153852|gb|AH000939.1|SEG\_STRTN916 [153852] gi|153851|gb|M22645.1|STRTN9162 [153851] gi|153850|gb|M20864.1|STRTN9161 [153850] gi|153660|gb|M36878.1|STRIF2BA [153660] gi|153585|gb|M13771.1|STRBRP [153585] gi|153575|gb|M64265.1|STRATPEFHA [153575] gi|153565|gb|M90060.1|STRATPASEA [153565] gi|152969|gb|M92376.1|STABLAIA [152969] gi|309660|gb|L14285.1|PCFPRGWZY [309660] gi|433714|gb|L12033.1|ENESATA [433714] gi|290645|gb|L15304.1|ENEVANB2A [290645] gi|148331|gb|M84146.1|ENEVANR [148331] gi|148329|gb|M64304.1|ENEVANH [148329] gi|148326|gb|M68910.1|ENEVANCRES [148326] gi|148324|gb|M75132.1|ENEVANC [148324] gi|148323|gb|L06138.1|ENEVANB [148323] gi|148321|gb|M85225.1|ENETETM [148321] gi|148320|gb|L00925.1|ENERTRNA [148320] gi|148319|gb|L00924.1|ENERRNA [148319] gi|148317|gb|M81466.1|ENERECA [148317] gi|148315|gb|M81961.1|ENENAPA [148315] gi|148312|gb|M38386.1|ENEMSPDPS [148312] gi|148310|gb|M37185.1|ENEGELE [148310] gi|148307|gb|L07892.1|ENEBLACREG [148307] gi|148305|gb|M60253.1|ENEBELAA [148305] gi|148303|gb|M77639.1|ENEB14NAM [148303] gi|290644|gb|L16515.1|ENERGTG [290644] gi|154954|gb|M37184.1|TRN916 [154954] gi|148301|gb|M69221.1|ENEAAD9A [148301] gi|148308|gb|M38052.1|ENECYLB [148308]



Table 28

# Phage Dp1 complete genome sequence. 56506 nucleotides.

1	ot 22t 2222	ratgaaggag	atattooott	aattattgct	taacaaaato	caccgaattt	atatataata	
						aggetegaac		
71								
141						caagacacca		
211						ataattacct		
281	gacggatatg	aattcacttc	cctttgccct	aaaacaggac	agcctgactt	cgcgaatgtt	ttcattagtt	
351	acattccaaa	cgaaaagatg	gttgaatcta	aatcattgaa	attgtactta	ttcagtttcc	gtaaccacgg	
421						tgatggaacc		
491						cgtcaacaaa		
						ttccttggaa		
561								
631						tatccggcgg		
701						tgctatagca		
771	gacaaaagca	tgaagcagaa	cttgaaaatg	ctgctaatgt	tgcaatgttc	tacggagtca	agttcaccat	
841	tcttgaaatt	gactcgaaaa	tctactcaag	ctctagctct	tccttattac	aaggaaaagg	cgaaatttca	
911	catqqaaaat	cttacgctga	aatcctagca	gagaaggaag	tagttgacac	ctatgttcca	tttagaaatg	
981						tacgtcgtat		
1051						ataattcaat		
1121						tctaaccaag		
1191		•		_		tatgaaagtg		
1261						atggaatgac		
1331	cattataagg	agaattgata	tgagagtttc	taaaacctta	acattcgacg	cagctcatca	actagttgga	
1401	cattttggaa	aatgcgcaaa	tttgcacggg	catacttaca	aagtcgaaat	ttcattagca	ggcggaactt	
1471	atgaccacgg	ttcqaqtcaa	gggatggttg	ttgactttta	tcacqtcaaq	aaaatcgcag	gtacattcat	
1541						tagcaaatgc		
1611						ccttacctgg		
1681						cctacaggtt		
1751						taacctttat		
1821	gaaaagatta	ctgtccgcga	aattttagag	caggagcagg	ataatggtta	atcaatacaa	tcagcctgaa	
1891	agaggcaaga	ttcgaatcaa	tgttcgcgac	cctgagaaaa	tgcctatcat	ggaaattttc	ggtcctacaa	
1961	ttcaaqqtqa	aggaatggtt	ataggtcaaa	agactatttt	cattcgaact	ggtggatgcg	actatcattq	
2031				-	_	atatcacagg	_	
2101						taaccacgtg		
2171						ctaaaagaac		
2241						taagcgatat		
2311	cctaaaccgc	cttcaagtgg	aatgagaact	aatatgaaaa	ttcttgaagc	tattgtagat	agaatgaatg	
2381	atgaaaacct	tgactggtca	tttaaaatcg	ttatctttga	cgaaaatgac	ctagcttatg	cgcgtgatat	
2451	gtttaaaact	ttcgaaggca	agttacgtcc	agtgaactac	ctttcagttg	ggaatgcaaa	cgcatacgaa	
2521	qaaqqaaaaa	tcaqtqataq	gcttcttgaa	aagttgggat	ggctttggga	taaagtgtat	qaaqacccaq	
2591						aataaaagag		
2661						ggatgggctt		
2731						ttggtctatt		
2801						cgctgaacat		
2871						gaagaccttg		
2941	agacattcca	ttcaattctt	tatgtgagca	tcatttagct	ccgttcgtag	ggaaggtgca	tattgcatac	
3011	attcctaagg	ataagattac	aggtctttca	aaattcggtc	gagtggttga	aggatacgct	aaacgacttc	
3081	aagtacaaga	gcgcttgact	caacaaatcq	ctgacgctat	tcaggaagtt	ctaaatcctc	aagcagttgc	
3151						agcacggggc		
3221						attgcttcag		
3291						agctcttatt		
3361						tttagagtca		
3431	•					gaacaattat	-	
3501	ttttcacctc	ttggactgat	tgacgtttta	ttcggttcac	ttgctacctt	ccttggagta	gtggcaatgg	
3571	tgaaagttgc	taagatggca	agtcctctat	attcacttat	ctgtccagtt	cttgctaatg	cttaccttat	
3641	tgcgctggaa	cttcgaatag	tttactcttt	acctttttqq	qaatctqtca	tctatgtagg	aattaqtqaa	
3711						caatcatttt		
3781						tagcactgac		
3851						attggcaaaa		
3921						gtgcatattc		
3991						cgtgggaatg		
4061	tegeegaact	cgataaaatt	cctggtgtat	ttagacagcc	taagacacgt	gaacagcttt	tggaagcacc	
4131	acaaatttct	tgggataatt	atctatacat	gcgcgagcga	atggttgaga	aagacaagct	cttacctatt	
4201						attcgaaggc		
4271						gacaagtgga		
4341						ttgggatgac		
		_	-		_		-	
4411						cacaggagcg		
4481						gatgctgtcc		
4551						attttagcct		
4621	gttgaggact	ataaacttcg	agcattgttc	aatgttcaat	acatgctgaa	ttgggcagag	aactatgaat	
4691	tcaagggaat	taaaaatcgt	caacgtcgac	tattttagat	aagagctttt	cgctcttatt	tttttaaaa	

. -

4761 aaaaatgaac tttttataca aaaacgcttg actttattca ctcattatcg tataatcata atataaataa 4831 aacqaataag aggtaaataa aatgacagca gttcaacaag ttaagttcta cttagaagaa gccggcqctc actttctaaa agatgttgag tacagtgaca acttagagca agcaattatg aaagatattc ttaaatggaa 4901 tqqcqctcat agagatgagc acgatatgaa aataacttca tacgaagtat tatagagagg ggtaaqqcta 4971 tgaaaaaagt tcaaacttat caagaatatc taaaactagt tgagttcaaa cgtcaacttt ctttaaatct 5041 tcgagaagga aaaataggag tcgatgaagc ggttattcaa ttattcacct tctatagttt caacaatatc 5111 gaggaacctc ctttcattgt actcaaaatg caagaggctg ccgtgaacgg gacttatgaa gcaaaactca 5181 5251 atatcaaaaa aaggaggctc atattatgag tattaagttc aaaaccgaag aactttcaaa aattgtttct 5321 5391 cageteaara agttgaagee tageaagttg ctagaaatea caaactattg geatattttt ggtgaeggeg aatqcqtcat qtttacagcg tatgatggct caaacttcct tcgatgcatt atcgacaqcg atqttqaaat 5461 tgacgtgatt gtgaaagcag agcagtttgg aaaacttgta gaaaagacca cggccgcaac cgtcacatta 5531 gttcctgaag aatcttcgct aaaagttatt gggaatggtg agtacaatat tgatattgtt acaqaagatg 5601 aagagtaccc tacattcgac cacttgctcg aagacgtgag tgaagaaaat gctctcactt tgaaaaagctc 5671 getgttetae ggaategeea atateaaega ttetgeggta tetaaateag gageagatgg aatttataee 5741 ggcttcctgt taaaaggcgg aaaagcaatt actacagaca tcattcgcgt atgtatcaac cctatcaagg 5811 aaaagggact agaaatgctc attoottaca acctaatgag tattttagca agtattootg atgagaagat 5881 5951 gractictgg casattgacg atactactgt ctatatttca toggottcag togaaattta toggaaaattg 6021 atggaaggta tggaagatta tgaagacgtt tcacagcttg actcaattga gtttgaagat gatgcggcta 6091 tecetacage agaaateetg agegtattag accecettgt actatteact teageetttg acaaaggaae cgtcgaattc ttattcttga aagaccgact tcgaattaaa acttctacta gcagttatga agacatcatg 6161 6231 tacgcatctg ctggcaagaa agtttcgaag aaagaattca cttgccacct taacagctta ctcttgaagg 6301 aaattgtatc aaccgtcacc gaagaaaact tcactgtctc ttatggaagc gaaaccgcaa ttaagatttc 6371 ategaatggt gtegtttact tectageact teaagageeg gaagaataat ggeeaagtee aatttaacta 6441 gaattgcaaa gatggttaga gcaggaaaca gtgaaggtcc tgcttcatct tttgtcaatt cgctgacccq ggttattgaa cgaactcagc ctgaatataa tccttcgaca tattataagc ccagcggggt tggtggatgt 6511 6581 attogaaaaa tgtattoga aagaatoggt gagtotatta tagataacgo agattotaac otaattqoaa 6651 tgggcgaagc tggaacattt aggcacgaag ttctccaaga gtacatggtt aaaatggctg aaatcgatga 6721 ggactttgaa tggttgaatg tagcagagtt cttgaaagaa aatccagttg aaggaactat cgtcgacgag 6791 cgtttcaaga aaaacgatta tgaaacgaag tgtaagaacg aacttcttca actttcattc ttgtgtgacg 6861 gactagtteg atataaagge aagetetaca ttttagagat taagaetgaa accatgttea agtteactaa acatactgag ccctatgaag aacacaagat gcaagcaact tgctacggaa tgtgtctagg agtcgatgat gtcattttcc tttatgaaaa tcgagataac ttcgaaaaga aagcctacac gtttcacatc acagacgaga 6931 7001 tgaaaaatca agtoottgga aaaattatga ootgogaaga gtatgtagag aaaggogaaa gtootaaaat 7071 ctattgctct tcagcctatt gcccatattg tagaaaggaa ggtcgaaatc tgtgagctat actggaaaaa 7141 tgttcgagga agactttttc gaaggtgcaa aagactttga gaaagatgct ttcacggtcc gtctatatga taccactaat ggattcgag gagttgcaaa tcctgcgat tatatagccg caactaactt tgggaccttg tttattgaac tgaaaactac taaagaagct tctttgagct ttaataacat cactgataat caatggttcc 7211 7281 7351 agctatcacg cgcagatgga tgcaaattta ttctcgccgg aattttagtg tatttccaaa agcatgaaaa 7421 gattatatgg tatccaattt caagccttga aaaaattaaa cggtctggag ttaaaagcgt caacccaaat ttcatcgatg cagggtatga agtttcttac aagaagcgtc gaactagatt gaccattcct ttccaaaatg 7491 7561 7631 ttctagatgc agttgagctt cattacaagg agaaaagcaa tggcaagacc taagttacct caaattgata ttcgagaaga agaaatacga gatgctcaag acgtagcaga ctcgtatggt gcgattatca ataaagtagt 7701 7771 cgacgaaatt gttgaagcag cttgcggttc acttgaccag gcaatggaag aaattcaaat agttgtaagc caaaatcctg tcattatgga agaccttaac tactacattg gctatcttcc cactcttctt tatttcgccg 7841 7911 cagatagggc ggaaatggtg ggaatacaaa tggattcaag ttctgctatc aggaaagaaa aatacgataa totatacatt tragcogoog ggaaaactat tootgacaag caagcagaaa ctogaaaact tgtcatgaat gaagaagtca tcgaaaatgc ttacaagcga gcctacaaga aagttcaatt aaagctagaa caggccgata 8051 aggtattagc atctttaaaa cgaattcaaa cctggcaact agcagagtta gaaactcagt caaataattc 8121 aaaaggagta ttattaaatg caaaaagacg tagacgtgaa aatgattgac cctaaacttg accgattaaa 8191 atacacaggt gattgggttg atgtacgaat tagttctatc actaaaattg acgccgacag cgccgatgtc 8261 8331 tcaagatgtc gaaaagtgct tcaaaaggct caagtatatt cagtggcggc aggtgaatgc attaaaattg 8401 cacacggatt tgctcttgaa cttcctaagg gatatgaagc aatcttgcat cctcgttcca gtctttttaa 8471 gaaaactggt ctaatcttcg tttctagcgg agtgattgac gaaggttaca aaggtgacac tgatgaatgg 8541 ttctcagttt ggtatgctac tcgtgacgca gatatcttct acgaccaaag aattgcccaa tttagaattc 8611 aggaaaagca acctgctatc aagttcaatt tcgtagaatc tttaggaaat gcggctcgtg gaggccatgg 8681 aagtacaggt gatttctaat gaaattggaa cagttgatga aggactggaa taaggattcg aaagctcttg 8751 tagcagttca aggacttgaa cgtgaagcgc ttccaagaat ccctttttct gcgccttcta tgaattatca 8821 aacctacggc gggctccctc gaaaaagggt agttgaattc ttcggtcctg agtcaagtgg gaaaactact tragetree acattetraa gaateegraa ateetatte agraegaate egaacagaag actegaagaac 8891 tcaaggaaaa gctggaaaat gggcgtgcat ccaaaggtag caagactgct gtcaaggaac ttgaaatgca actcgatagt cttcaagagc ctcttaagat tgtatatctt gaccttgaga atacattaga cactgagtgg 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354

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accttatega caagetaaaa gaggageate tatataatte acttgtteea attttaaegg aageggetga ggacattcaa gtagatagta acattgcgat tgcgaatata attccaaaac tagaagaact tttcaatcgc 49911 tctaaattcg taggcggact agacattgct cgaaatgcta aacttcgact agactgggcg aatactatta 49981 50051 gaaaccatga cggtgaaaga cttggaatat cgacagggtt tgaactattg gacgacgtgc ttggaggctt gadattatga taggatttga tigtcataat ggctcgacct ggacaaggta agtcgtggac tattgataaa atgcttgcaa ctgcttggaa gaacgggcat gatgtccttc tatatagcgg ggaaatgagt gaaatgcaag 50121 50191 trggtgctcg tatagatact attotttcga atgttagcat caattcaatt accaaaggga trtggaacga 50261 ccatcagttc gaaaaatatg aggaccatat tcaagcaatg actgaggctg aaaattccct tgtggtagtc 50331 50401 acgcccttta tgattggagg aaagaacctt acccctgcaa ttttagatag catgatatct aaatatagac 50471 catctgtggt ggggattgac cagctttcac tcatgagcga gtcttatcca agcagggagc agaagcgaat ccagtacgcc aacatcacca tggacctata taagatttct gctaaatatg gaattcctat tgtgcttaat 50541 gtccaagcag ggcgttcggc taaaactgaa ggcgctgaaa gtatggaact agaacatata gcagaaagtg 50611 atggagtagg tcaaaatgct agcagagtta tcgctatgaa gcgtgacgaa aaatccggca tacttgaact 50681 50751 atctgtcgtt aaaaaccgat atggcgaaga ccgaaaaatc atcgaatata tgtgggacgt tgaaactgga 50821 acctatactc ttataggatt caaagaggaa ggcgaagaag gaactgaaaa aggcgaaagc tctccattga 50891 aagcaaaagc ctctaggtcg actgctcgtc ttcgaagtaa ggttacaagg gaaggagttg aagcattttg 50961 atgaaagtaa atggtottoa aattgaagog actootgaac aaataattga aaaactttog agacaacttg 51031 aagacgaagg aacattcatt tttagacgaa ctaagtcgct tggaagcaac tatcaattct catgcccqtt 51101 tcatgcagga gggactgaaa agcatccctc ttgtggcatg agtagaaatc cttcttattc aggaagtaag 51171 gtgacggaag ctggaacggt tcactgtttc acttgcggct acacttcagg actaactgaa ttcgtctcga 51241 atgtattagg tcgaaacgat ggagggttct atggaaacca gtggctgaaa aggaattttg gaacatctag 51311 cgaagtagtt aggcaaggcg tcagccctga agcgtttcga agaaatggga gaactgaaaa agtcgagcat 51381 aaaatcattc ctgaagagga acttgataaa taccggttta ttcatcctta tatgtatgaa cggaaattga 51451 eggacgaget categagatg titigatgtag gitatgacaa actgeatgat tgeateacet titecagtaeg gaacctcaag ggcgaaacag tattcttcaa ccgtcgaagt gttcgttcta agtttcacca gtacggtgaa 51521 gatgacccta aaacggaatt tctttatggc caatatgagc ttgtagcatt tcgagactat tttgaaaaac 51591 ctattagtca agtattcgtg actgagtctg ttatcaactg cttgactctt tggtcaatga agattccagc 51661 agtocotott atgggagtag gtggaggaaa toaaatoaat ttactaaaac gacttootta tagaaatatt gttotagoac ttgacootga taacgotggg cagacagog aggaaaaact ctacogacag ttaaagogaa 51731 51801 gcaaggtcgt tagatttttg aactacccta aagagttcta tgataataag tgggatataa acgaccatcc ggaattatta aattttaatg attragtctt gtagaaattc atttattatc gtataataaa gtagaaaat 51871 51941 tttaaaaaga ggtcatatca atatgaaaga agcgaataga ctagtttcta gctatgtagg attcgaatgc 52011 tggactgacg aagaatgtat caggaacttt gaactagace etgatatgte aattgegtet gettateate 52081 52151 gttattttgg gatgctttat tcctatgcaa aaaggtttaa atgcttatct cgacatgaca ttgaaagcat tgcattcgag actatttcaa aatgtttggc aacgttcaaa tcaaaccaag gggccaagtt ttcaaactac cttacaagac tcttcaagaa tagaatagtc ttagaatata ggtacctaaa tgcaccttcc atgaatcgaa 52221 52291 52361 attggtatgt agaagtgacg ttcgatagcg tttcgacaaa tgaagaaggc gacgatttta gtatcctatc 52431 gacagttggc tattgtgaag actacggaaa aattgaaatt gaagcaagtc ttgacttcat gacgctttct 52501 aatacagagt atgettatat ctegtetgte atteaaaaeg gteetteagt aagegaegea gaaattgege 52571 gtgaaattgg agtaagcagg tctgctatta gtcagtctaa gaagtcacta aaaaataaat taaaagattt

atteatteat tattat

tatataactg gtttacaaat cacgtgaatt tcgtgtatat tatatatgaa aggacaaact ttgaaacctt 52641 aaaaacttca aaaatctttc aaccattaaa aacttataaa ggagaatcga tatgggaaaa gtatcaattc aaaaatcagg aacatttagc tcagggtcta ataacgagtt tttcacactc gctgaccacg gtgacagcgc 52711 52781 52851 aattgtcact ctattgtatg atgacccgga aggcgaagac atggattatt tcgtagtcca cgaagcagac gttgacggtc gtcgacgcta tatcaattgc aatgctattg gcgaagacgg ggaaacagtc catcctgata 52921 52991 attgtccatt atgccaaaac ggattccctc gtattgaaaa actatttctt caactttaca accatgatac 53061 gggaaaagtt gaaacatggg accgaggccg ttcttatgtt caaaagattg ttacatttat caataaatat 53131 ggaageettg tgaeteagee ttttgaaatt attegtteag gagetaaagg tgaecaaega actaettatg 53201 aatteettee agagegteeg gaagacagtg etactettga agatttteea gaaaagageg aacttettgg aactctaatt ttagacctcg acgaagacca aatgtttgac gtggttgacg gcaagttcac tcttcaagaa gagcgttctt caagtcgttc aaattcacgt agaggagcat ctcctgcgcc tagacgaggt tccggtcgag 53271 53341 53411 aatottoaca aggtogaaca gotgaaagaa otoottoagt tagtogaaga actootocaa cacgaggtog 53481 aggattctaa catgagggcg cgagccctct ttattattga ttaagaaagg gaaaataatg gcacaaaaag gactetttgg tgcaaageet egttetagea agaagaaega tgeteagtta ettgeteaae ggaaaaaeag 53551 gaageetgea gttgaggtta ettacattte aggaaaeget etaaaggaeg eagttgetag agetegtaet 53621 ctttcaacta ggattettgg acacgttett gatagaettg agttaateae tgaggaagea aaactegage 53691 agtatgtaga caaaatgatt gaagacggaa taggttctat tgacgtagaa actgatggac tcgatactat 53761 tcacgatgag ctggcaggag tctgcttgta ctcacctagt caaaaaggaa tctatgctcc tgtcaatcat 53831 53901 gttagcaata tgacgaagat gcgaattaag aatcaaattt ctcctgagtt catgaagaaa atgcttcaac ggattgtaga ttcaggaatt cotgtoatot atcataattc gaaatttgac atgaaatcga tttattggcg 53971 actoggogto aaaatgaatg agcoagogtg ggatacatat ttagoogcaa tgottttaaa tgaaaacgag 54041 54111 teteacaget tgaaaagtet teactetaaa tatgttagga acgaagaaaa egeagaggtt geaaaattta 54181 atgacttatt taaaggaatt cettttagtt taatteetee tgatgttgee tatatgtatg eggeetatga 54251 ccctttgcaa actttcgaac tctatgaatt tcaagaacaa tacttgactc caggaactga acaatgtgaa 54321 gaatataacc tggaaaaagt ctcatgggtt cttcataata ttgagatgcc tctaattaaa gttctcttcg 54391 acatggaagt ctacggtgtc gacttagacc aagataagct ggcagaaatt agagaacagt ttactgccaa 54461 tatgaacgag gctgagcaag agtttcaaca gcttgtcagc gaatggcagc ctgaaattga agaacttcga 54531 canactaatt tocagagota toaaaaacto gaaatggatg caagaggtog agtgacggta agcatttoca 54601 gtcctactca attagcaatt ctgttttatg atatcatggg attgaaaagt cctgaaaggg ataaacctag aggaacaggc gaaagtattg tcgagcattt tgataacgat atctcaaaag cacttttgaa atatagaaaa 54671 tatgcaaaat tagtttcgac ctatacaaca cttgaccaac accttgcaaa gcctgacaat cgaattcaca 54741 54811 ctacattcaa acagtacgga gctaagacag ggcgtatgtc aagtgagaat cctaacttac agaatattcc 54881 ttctcgcggt gagggtgcag tagttcgaca aatctttgca gccagtgaag ggcattacat tattggtagt 54951 gactactete aacaagaace tegtteattg geggaattaa gtggegaega aagtatgega eatgettaeg 55021 aacaaaacct ggacctatat tcagttatcg gttcgaaact ttatggtgtt ccctatgaag agtgtttaga 55091 gttctatccc gacggaacga ctaacaagga aggaaaactt cgaagaaatt ctgtcaagtc cgttctttta 55161 ggtcttatgt acggccgcgg ggctaactca atcgctgagc agatgaatgt atctgtcaaa gaagcgaata 55231 aggttattga agatttette accgagttee etaaagtgge agactatate atattegtte aacagcagge 55301 graggarttg ggatatgtte aaacagetae eggtegaaga agaaggette etgatatgag tetteetgaa 55371 tacgagttcg agtatatcga cgctagcaag aacgaagatt tcgacccctt taactttgac gcagaccaac 55441 agatggacga tactgttcct gaacatatta tcgaaaaata ttgggcccag ctagatagag cctggggatt 55511 taagaagaag caagaaatta aagaccagge aaaagccgaa ggaattetta ttaaggataa cggaggcaag 55581 atagetgatg etcagegeea atgettgaac teagetatte aaggaacgge ageegacatg actaagtacg 55651 caatgattaa ggtacacaat gacgctgaat tgaaagaatt aggattccat ttaatgattc cagttcacga tgagttacta ggtgaggttc ctatcaagaa cgcaaaacgg ggagcagaaa ggttgacaga agttatgatt gaagcagcca aggacattat tagtcttcca atgaaatgtg accccagtat agtagaaaga tggtatggtg 55721 55791 aagaaattga aatctaaaat ctattcagtt gcatatataa ttctagtagt tattgcgaac cttgtgacaa 55861 tttatttcga acctttaaat gtgaaaggaa ttttaattcc tccaagcagt tggtttatgg gattcacttt 55931 cctgcttata aatctaataa gcaagtacga gaagccaaaa tttgcaggtt ctttgatatg ggtagggtta 56001 56071 tteettacet egitgatitg etitatgeaa aacetaceae aategeitgt egitggeitea ggagtigeat 56141 tttggataag tcaaaaagca agtgtcttta tattcgacaa gctctcgaat aaattagact cgaagattgc aaatgetttg tetageaaca teggttetat tatagaegea accatatgga ttteattagg actgagteet 56211 56281 cttggaattg gaacggttgc atatatagat attccgtcag ccgtactagg ccaagttcta gttcagttta 56351 tottgcagtc aattgcttcg agatatttga aaaagtagtc aggaaaattc ctgattatct tgcaqtcaat 56421 tgcttcgaga tatttgaaaa agtagtcagg aaaattcctg attattttt ttacaaaaac gcttgacttt

Table 29

# Phage dp1 ORFs list

up	Name	Frame	Position	Size (a.a.)	Key words
1	dp1ORF001	2	3669840390	1230	Putative tail;
2	dp1ORF002	1	3238635835	1149	Tail;
3	dp1ORF003	3	5353855877	779	DNA polymerase I;
4	dp1ORF004	3	4040142440	679	Minor structural;
5	dp1ORF005	1	2367425434	586	
6	dp1ORF006	2	4529646987	563	SWI/SNF Helicase;
7	dp1ORF007	3	2223023621	463	Terminase;
8	dp10RF008	1	4962450961	445	DNAb Helicase;
9	dp1ORF009	2	1316014404	414	
10	dp1ORF010	2	86999859	386	RecA;
11	dp1ORF011	3	2801729096	359	Major head;
12	dp1ORF012	3	53466419	357	DNA pol. III beta;
13	dp1ORF013	3	1021511240	341	DNA pol. III gamma and tau;
14	dp1ORF014	3	5096151974	337	DNA primase;
15	dp1ORF015	1	37934728	311	
16	dp1ORF016	3	4341344303	296	Amidase:
17	dp1ORF017	1	1124212081	279	
18	dp1ORF018	3	3584736686	279	
19	dp10RF019	2	1216112967	268	
20	dp1ORF020	1	18642658	264	exsD; Coenzyme PQQ;
21	dp1ORF021	2	25043295	263	GTP cyclohydrolase;
22	dp1ORF022	2	3089631675	259	
23	dp1ORF023	2	64197195	258	
24	dp1ORF025	-1	1802618778	250	
25	dp1ORF024	3	2599226738	248	
26	dp1ORF026	2	2151222252	246	
27	dp1ORF027	1	5276253490	242	
28	dp1ORF028	3	4459545299	234	
29	dp10RF029	2	6621348	228	exsB;
30	dp1ORF031	3	2694327611	222	6,30,
31	dp1ORF030	-2	1942320088	221	
32		1	5203352647	204	
33	dp1ORF032 dp1ORF033	2	76708239	189	
34	dp1ORF035	-1	1685917425	188	
35	dp1ORF036	1	4880849362	184	DNAc replication;
36	dp10RF037	1	5585556388	177	DIVAC (epitcation,
37		2	131652	173	
38	dp10RF034	3	13501871	173	ovoC: 6 pump audintenhadoratoria.
	dp10RF038	3	33063803	165	exsC; 6-pyruvoyltetrahydropterin;
39	dp1ORF039			_	Citrulline biosynthesis:
40	dp1ORF040	1	71927683	163	4070
41	dp1ORF041	3	82088699	163	dUTPase;
42	dp1ORF042		4808248561	159	
43	dp1ORF043	1	3169932154	151	
44	dp1ORF044	-1	2521125666	151	
45	dp10RF045	2	2534025777	145	
46	dp1ORF046	3	4277443202	142	
47	dp10RF047	1	4754247961	139	
48	dp1ORF048	-3	1630816709	133	
49	dp1ORF049	-3	4362044018	132	
50	dp1ORF050	3	1508115476	131	
51	dp1ORF051	2	2976530154	129	
52	dp1ORF053	-3	4991750300	127	
53	dp1ORF052	3	3051630893	125	
54	dp1ORF054	2	1442314800	125	
55	dp1ORF055	3	2762728004	125	
56	dp1ORF056	-3	1878019151	123	* **
57	dp1ORF057	1	985910218	119	-
58	dp1ORF058	3	1563315989	118	
	dp1ORF059	1	3015430507	117	
59					
59 60	dp1ORF060	-2	3771738070	117	
60 61	dp1ORF060 dp1ORF062	-2 -3	4494045284	114	
60	dp1ORF060				

64					360	
65	64	dp1ORF066	-3	2856628898	110	
67					108	
B	66	dp1ORF068	3	2945129768	105	
B9	67	dp1ORF069	-3	2009420411	105	
101 dp10RP072 - 2 50749,51045 98 9	68	dp1ORF061	-3	1916119475	104	
71 dp10RF072 -2 50749,51045 98 97 72 dp10RF073 3 14262,14555 97 7 7 7 dp10RF075 -1 22154,22447 97 7 7 dp10RF075 -1 52154,22447 97 7 7 dp10RF076 -1 5435,5728 97 7 6 dp10RF076 -1 5435,5728 97 7 6 dp10RF076 -1 5435,5728 97 7 6 dp10RF077 -1 14800,15084 94 94 94 94 94 94 94 94 94 94 94 94 94	69	dp1ORF070	1	1597316284	103	
72 dp10RP074 3 32298,3291 97 73 dp10RP074 3 32298,3291 97 75 dp10RP076 -1 22154,22447 97 75 dp10RP076 -1 45435,7528 97 76 dp10RP076 -1 14800,15084 94 77 dp10RP079 -3 35007,33288 93 78 dp10RP079 -3 35007,33288 93 78 dp10RP03 -3 49382,48527 91 80 dp10RP03 -1 44208,42759 89 81 dp10RR03 -1 44728,44994 88 82 dp10RR03 -1 44728,44994 88 82 dp10RR08 -1 44728,44994 88 83 dp10RR08 -1 35728,3374 84 84 dp10RR08 -1 35728,3374 84 85 dp10RR08 -3 10602,10847 81 86 dp10RR08 -3 10602,10847 81 86 dp10RR08 -3 10602,10847 81 87 dp10RR08 -3 10602,10847 81 88 dp10RR08 -3 10602,10847 81 89 dp10RR08 -3 10602,10847 81 89 dp10RR08 -3 10602,10847 81 80 dp10RR08 -3 38883,1910 70 90 dp10RR08 -1 4689,46881 70 90 dp10RR08 -1 4689,46881 70 90 dp10RR08 -1 48692,46881 70 90 dp10RR08 -1 48692,46881 70 90 dp10RR08 -1 48692,46881 70 90 dp10RR09 -1 38883,1910 70 90 dp10RR09 -1 1597,1803 68 90 dp10RR09 -1 1597,	70	dp1ORF071		3890439209	101	
73 dp10RP075 -1 22154,2247 97 74 dp10RP075 -1 52154,2247 97 75 dp10RP076 -1 5435,5728 97 76 dp10RP077 -1 14800,18084 94 77 dp10RP079 -3 35007,33288 93 78 dp10RP081 -3 55188,55466 92 79 dp10RP103 2 49352,49627 91 80 dp10RP080 1 42490,42759 89 80 dp10RP080 1 44728,44994 88 80 dp10RP080 1 44728,44994 88 81 dp10RP080 1 44728,44994 88 82 dp10RP080 -3 51264,51497 83 83 dp10RP085 -3 16602,18047 81 85 dp10RP087 -2 29794,30036 80 86 dp10RP087 -2 22794,30036 80 86 dp10RP087 -2 12256,12495 79 87 dp10RP087 -2 12256,12495 79 88 dp10RP087 -3 17280,17507 75 90 dp10RP078 -3 17280,17507 75 90 dp10RP091 1 43188,43413 74 91 dp10RP090 1 43188,43413 74 92 dp10RP090 1 43188,43413 74 93 dp10RP090 1 43188,43413 74 94 dp10RP090 1 43188,43413 74 95 dp10RP090 1 43888,39100 70 96 dp10RP090 1 43888,39100 70 97 dp10RP090 1 43888,39100 70 97 dp10RP090 1 1 5220,14266 69 98 dp10RP090 1 43888,39100 70 99 dp10RP090 1 1 5220,1426 69 99 dp10RP090 1 43888,39100 70 97 dp10RP090 1 43888,39100 70 97 dp10RP090 1 43888,39100 70 97 dp10RP090 1 1 5220,1426 69 98 dp10RP090 1 1 5220,1426 69 99 dp10RP090 1 1 5220,1426 69 90 dp10RP090 1 1 5220,1426 69 90 dp10RP090 1 1 5220,1426 69 91 dp10RP090 1 1 5220,1426 69 91 dp10RP090 1 1 5220,1426 69 91 dp10RP090 1 1 5220,1426 69 91 dp10RP090 1 1 5220,1426 69 91 dp10RP090 1 1 5220,1426 69 91 dp10RP100 1 1597,1603 68 91 dp10RP100 1 1597,1603 68 91 dp10RP100 1 1597,1603 69 91 dp10RP100 1 1597,1603 69 91 dp10RP100 1 1597,1603 69 91 dp10RP100 1 1597,1603 69 91 dp10RP100 2 2 31435,31830 69 91 dp10RP100 1 1597,1603 69 91 dp10RP100 2 2 31435,31830 69 91 dp10RP100 1 1597,1603 69 91 dp10RP100 1 1597,1603 69 91 dp10RP100 1 1597,1603 69 91 dp10RP100 1 1597,1603 69 91 dp10RP100 1 1597,1603 69 91 dp10RP100 1 1597,1603 69 91 dp10RP100 1 1597,1603 69 91 dp10RP100 1 1597,1603 69 91 dp10RP100 1 1597,1603 69 91 dp10RP100 1 1597,1603 69 91 dp10RP100 1 1597,1603 69 91 dp10RP100 1 1597,1603 69 91 dp10RP100 1 1597,1603 69 91 dp10RP100 1 1597,1603 69 91 dp10RP100 1 1597,1603 69 91 dp10RP100 1 1 1597,1603 69 91 dp10RP100 1 1 1597,1603 69 91 dp	71	dp1ORF072	-2	5074951045	98	
T4	72	dp1ORF073	3	1426214555	97	
15	73	dp1ORF074	3	3229832591	97	
76	74	dp1ORF075	-1	2215422447	97	
77 dg10RF079 3 35007.35288 93 78 dg10RF081 3 55188.55466 92 79 dg10RF03 2 48352.49527 91 80 dg10RF03 1 42490.42759 89 81 dg10RF03 1 4478.44994 88 82 dg10RF03 1 35720.35974 84 83 dg10RF03 3 51246.51497 83 84 dg10RF03 3 51246.51497 83 84 dg10RF03 3 51246.51497 83 85 dg10RF03 1 3 10602.10847 81 86 dg10RF03 2 29794.30036 80 86 dg10RF03 2 29794.30036 80 86 dg10RF03 2 29794.30036 80 86 dg10RF03 3 500.5279 79 87 dg10RF03 3 56256.55498 78 89 dg10RF03 3 1226.17250 77 89 dg10RF03 1 2703.72761 74 89 dg10RF03 1 2703.72761 74 91 dg10RF03 1 2703.72761 74 91 dg10RF03 1 43189.43413 74 Holin; 92 dg10RF03 2 45538.45756 72 93 dg10RF03 2 45898 70 95 dg10RF03 3 8878.9897 70 95 dg10RF03 3 8878.9898 70 95 dg10RF03 3 8878.9898 70 95 dg10RF03 1 43888.39100 70 96 dg10RF03 1 38888.39100 70 97 dg10RF03 1 43627.43536 89 98 dg10RF03 1 43628.4884 87 99 dg10RF03 1 43628.4884 89 99 dg10RF03 1 43628.4888 89 90 dg10RF03 1 43628.4888 89 90 dg10RF03 1 43628.4888 89 91 dg10RF03	75	dp1ORF076	1	54355728	97	
78	76	dp1ORF077	1			
	77					
80	78	dp1ORF081				
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241	dp1ORF240	1	4189342000	35	
242	dp10RF241	-1	4691347020	35	
243	dp1ORF242	-1	4123141338	35	<u> </u>
244	dp1ORF243	-2	5119951306	35	
245	dp1ORF244	-3	2697627083	35	
246	dp1ORF245	-3	61716278	35	· · · · · · · · · · · · · · · · · · ·
247	dp1ORF246	-3	27242831	35	
248	dp1ORF247	1	2964129745	34	
249	dp1ORF248	1	5356053664	34	,
250	dp1ORF249	2	20122116	34	
			2383723941	34	
251	dp1ORF250	2			
252	dp1ORF251	-1	3910139205	34	
253	dp1ORF252	-2	5466754771	34	
254	dp1ORF253	-3	5615156255	34	
255	dp1ORF254	-3	4837548479	34	······································
256	dp1ORF255	-3	94689572	34	
257	dp1ORF256	1	1528915390	33	
258	dp10RF257	1	2821628317	33	
259	dp1ORF258	1	4402344124	33	
260	dp1ORF259	2	42984399	33	
261	dp10RF260	2	2474624847	33	
262	dp10RF261	3	288389	33	
263	dp1ORF262	3	94089509	33	
			2695127052	33	· · · · · · · · · · · · · · · · · · ·
264	dp1ORF263	-1			
265	dp1ORF264	-1	60386139	33	
266	dp1ORF265	-1	47004801	33	
267	dp1ORF266	-2	5011950220	33	
		-2	4726647367	33	
		-/	4120041301		
268	dp1ORF267		40500 40004		
269	dp1ORF268	-2	1252012621	33	
			1252012621 5373353834	33	
269 270	dp1ORF268 dp1ORF269	-2 -3	5373353834	33	
269 270 271	dp10RF268 dp10RF269 dp10RF270	-2 -3 -3	5373353834 5069150792	33 33	
269 270	dp1ORF268 dp1ORF269	-2 -3	5373353834	33	

## Table 30

## Predicted Dp-1 amino acid sequences

dp10RF001 36698 atgattgacaataatttacctatgagtccaattcctggcgaaattgttcaagtatatgaccaaaacttcaatctaattggagca M I D N N L P M S P I P G E I V Q V Y D Q N F N L I G A 36782 agtgatgaaatctttagcaagcattacgaagacgaaattgtgactcgagctcgaggaaaagaaactttcacttttgaaagtatt S D E I F S K H Y E D E I V T R A R G K E T F T F E S I 36866 gaaacctcatctatctatcaacacttaaaggttgaaaacattatccagtatggaggaagatggtttcgaattaaatatgctcag ETSSIYQHLKVENIIQYGGRWFRIKYAQ 36950 gacgtagaagatgtcaaagggcttaccaagtttacctgctacgcattatggtatgaactagcagaaggcttgcctaggaagttg E D V K G L T K F T C Y A L W Y E L A E G L P R K L 85 37034 aaacacgttgcttcttctgtaggcgctgtcgcgctagatattatcaaagacgcaggtgaatgggttcgactagtttgtcctcct113 K H V A S S V G A V A L D I I K D A G E W V R L V C P P 37118  ${\tt gacggtgctaacaacaagttcgaagcataacagccgcagaaaattcaatgcttttggcatcttcgatatctttgcaaagcaatac}$ 141 D G A N K Q V R S I T A A B N S M L W H L R Y L A K Q Y 37202 169 N L E L T F G Y E E I I K Q E V R I V Q T V V F L Q P Y 37286 gtcgagtctaaagtagactttcctcttgtagttgaagagaatttgaaatatgtcactaggcaggaagattctcgaaacctgtgt 197 V E S K V D F P L V V E E N L K Y V T R Q E D S R N L C 37370  ${\tt acggcttacaagttgacaggtaaaaaggaaggaagtcaagagcctttaacgtttgcttctatcaacaatggaagtgaatat}$ TAYKLTGKKEEGSQEPLTFASINNGSEY 225 37454  $\verb|ctcattgatgtttcgtggtttactacacgccacatgaagcctcgatatattgctaaatctaaaagcgacgaacattttagaatt|$ 253 LIDVSWFTTRHMKPRYIAKSKSDEHFRI aaagaaaatttgatgagtgctgcgcgtgcttatcttgacatctacagtcgcccactaattggatatgaggcttcagcggtcctt K E N L M S A A R A Y L D I Y S R P L I G Y E A S A V L 37538 281 37622 tataacaaggttcctgacttgcatcatactcaactaattgtcgacgaccattatgatgttatcgagtggcgaaagatatctgct PDLHHTQLIVDDHYDVIEWRKISA 309 cgaaaaattgactacgacgacctttcaaactctactatcattttccaagaccctcgaaaagacttgatggacttgctaaatgag 37706 RKIDYDDLSNSTIIFQDPRKDLMDLLNE 337 37790  $\tt gacggcgaaggagtcctttcaggggaaactgtaaatgagtcccaagttgttattagatacgcagatgacattttagggactaat$ D G E G V L S G E T V N E S Q V V I R Y A D D I L G 365 tttaatgcagaatctgggaaatacattggtgtccttaatactaataagaaaccgagcgaattagttcctgacgactttacatggFNAESGKYIGVLNTNKKPSELVPDDFTW 37874 393 37958 attegactagaaggteetaaaggtgaegeaggtttacegggageteetgggegtgatggagtegaeggtgtacetggaaagage I R L B G P K G D A G L P G A P G R D G V D G V P G K S 421 38042 V G I A D T A I T Y A V S V S G T Q E P E N G W S E Q 449 38126 gttcctgaactcataaaaggtcgattcttgtggactaaaacattttggagatatactgacggctcacatgaaactggatactcc 477 PELIKGRFLWTKTPWRYTDGSHETGYS 38210 505 AYIGQDGNSGKDGIAGKDGVGIAATE 38294 atgtatgcaagttcgccatctgctactgaagctccagctggtggtctacgcaagttcctaccgtcccaggtggtcagtat M Y A S S P S A T E A P A G G W S T Q V P T V P G G Q Y 533 38378 ttatggactcgaacaagatggcgctacactgaccaaactgatgaaattggatattcagtttcaagaatgggcgagcagggtcct 561 L W T R T R W R Y T D Q T D E I G Y S V S R M G E Q 38462  ${\tt aaaggtgacgcaggtcgtgacggtattgcaggaaagaacggaatagggttgaagtcaacttcagtttcttatggaattagtccc}$ 589 K G D A G R D G I A G K N G I G L K S T S V S Y G I S 38546 actgattctgcgattcctggagtatgggcttcacaagttccttctttaatcaaaggtcaatatctttggactcgaactatttgg 617 T D S A I P G V W A S Q V P S L I K G Q Y L W T R T I 38630 645 38714 ggtaaggatggggtaggaattaagtctacgaccattacctacgcaggctcaacctcaggaacagttgcgcctacttcaaattgg G K D G V G I K S T T I T Y A G S T S G T V A P T S N W 673 acttctgctattccaaatgttcaaccgggattcttcttgtggacgaaaactgtttggaactatactgatgacactagcgaaaca T S A I P N V Q P G F F L W T K T V W N Y T D D T S B T 38798 701 38882  $\tt ggttactcagtttccaagataggtgaaacaggtcctagaggagttcaaggtcttcaaggtcctcaagggcttcaaggaattcct$ Y S V S K I G E T G P R G V Q G L Q G P Q G L Q G I 729 ggacctgcaggagctgacggacgttcgcaatatactcacctcgctttctctaatagtccaaacggtgagggatttagtcatact 38966 G P A G A D G R S Q Y T H L A F S N S P N G E G F S H T 39050 gacagcggacgagcatacgtcggtcagtatcaagatttcaatcccgtccattcaaaagaccctgcagcctatacatggacgaaa D S G R A Y V G Q Y Q D F N P V H S K D P A A Y T W T K 39134 tggaaggggaatgacggagctcaagggatacccgggaagccaggcgcagacggtaagactaattatttccatatagcttacgct 813 W K G N D G A Q G I P G K P G A D G K T N Y F H I A Y A 39218 S S A D G S R E F S L E D N N Q Q Y M G Y Y S D Y E 841 39302 gatagcagggatcgaactaagtatcgatggtttgaccgccttgccaatgttcaagtgggaggtcgaaacgagttccttaattct 869 D S R D R T K Y R W F D R L A N V Q V G G R N E F L N S 39386 897 EFGLKPRYSSYNLMDGQDQTQGQISA 39470 925 T I D E R Q R F K G A N S L R L D S T W N G K P Q N Q K

32470

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32722

32806 141

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169 32974

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33058 225

33142

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33730 449

33814 477 33898

505 33982

533 34066

561

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673 34486

701

34570 729

34150

34234

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34402

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57 32638 85

39554 ctgaccttttctttaggaggagatacgcgattaggtactccaaccgagtggtctaatttagaaggtcgtatcagtttctgggct 953 L T F S L G G D T R L G T P T E W S N L E G R I S F W A 39638 aaggcctctaggaacggagtgagcttagctgcacggccgggttatcgtagtaacgtatttaccgcaaccttaaccgatcaatgg 981 K A S R N G V S L A A R P G Y R S N V F T A T L T D Q 39722 aagttctacgattttaaattctttgacaaagttaattcaaattgtaccgctgaagcaattttccatgtattcactcaaagttgt 1009 K F Y D F K F F D K V N S N C T A E A I F H V F T Q S C 39806 1037 S V W L N H I K I E L G N I S T P F S E A E E D L K Y R 39890 attgactcaaaagccgatcaaaagctaactaaccaacagttgacggcactcacggaaaaggctcaactacatgacgcagaactg I D S K A D Q K L T N Q Q L T A L T E K A Q L H D A E L 1065 39974 aaagctaaggctacaatggagcagttaagtaacttagaaaaggcttatgaaggtagaatgaaagctaatgaagaagctatcaaa K A K A T M E Q L S N L E K A Y E G R M K A N E B A I K 1093 40058 aaatcggaagccgacctaatcttagcggcaagtcgaattgaagctactatccaagaacttggcgggctacggggaactgaagaag 1121 K S E A D L I L A A S R I E A T I Q E L G G L R E L K K 40142 ttcgtcgacagttacatgaggtcttattgaaggtctaattatcggtaagaacgacggtagctctaccattaaggtatcaagt 1149 F V D S Y M S S S N E G L I I G K N D G S S T I K V S S 40226 1177 40310 40390

1205 Q S I Q V G R F R T E Q Y S F N P D M N V I R Y V G \* dp10RF002

atggatttttgggtcaattgcagcaaaaatgactttggatatctcaaacttcacaagtcaattaaatcttgctcaaagtcaagcg M D F G S I A A K M T L D I S N F T S Q L N L A Q S Q A  ${\tt caacggctcgcactagagtcttcgaagtcctttcaaattggttctgctttaacaggattagggaaaggacttacgactgcggtt}$ Q R L A L E S S K S F Q I G S A L T G L G K G L T T A V gcggctcaaggtatggaaaatctagcttcagccggtttccaggtaaatgaaatcatggacgctatgccaggggtacttgacctg A A Q G M E N L A S A G F Q V N E I M D A M P G V L D L gctgccgtatctggaggagatgtggccgcgagctccgaggccatggctagttcacttcgagcctttggattagaggcaaaccag A A V S G G D V A A S S E A M A S S L R A F G L E A N Q gcgggtcacgtggctgacgtatttgctcgagcagcagctgatacgaacgcagaaactagcgacatggcagaggcgatgaaatac G H V A D V F A R A A A D T N A B T S D M A E A M K Y gtcgcacccgttgctcactctatgggcttgagccttgaagaaacggctgcgtctattgggattatggccgacgccggtattaag V A P V A H S M G L S L E E T A A S I G I M A D A G I K ggctcgcaagccggaaccacgcttagaggcgctctctcgcgtattgccaaacctacgaaagcgatggtcaaatcaatgcaggaa G S Q A G T T L R G A L S R I A K P T K A M V K S M ttaggagtttcgttctacgacgcgaacggaaacatgattccactaagagaacaaatcgctcaactgaaaacagctactgcagga L G V S F Y D A N G N M I P L R E Q I A Q L K T A T A G ctaacacaagaggaacgaaatcgtcaccttgttaccttgtatggccaaaactcgttgtcaggtatgcttgcactattagacgcaLTQBERNRHLVTLYGQNSLSGMLALLDA G P E K L D K M T N A L V N S D G A A K E M A E T M Q D aaccttgctagtaaaatcgagcaaatgggaggagctttcgagtctgttgctattattgttcaacaaatccttgagcctgcactt N L A S K I E Q M G G A F E S V A I I V Q Q I L E P A L gctaaaatcgtgggagcaatcacaaaagttctcgaagcattcgtaaatatgtcacctatcggtcaaaagatggttgtcatattc A K I V G A I T K V L E A F V N M S P I G Q K M V V I F gcaggaatggttgcagcccttggaccactgcttctaattgcaggaatggtgatgacaactattgtcaagttaagaattgctatt A G M V A A L G P L L L I A G M V M T T I V K L R I A I  ${\tt cagtttttaggtccagcatttatgggaacgatgggaaccattgcaggagttatagcaatattctatgctctggtcgccgtgttc}$ Q F L G P A F M G T M G T I A G V I A I F Y A L V A V atgatagcctacacaaaatcggagagatttagaaactttatcaacagtcttgcgcctgctattaaagctgggtttggaggagcg M I A Y T K S E R F R N F I N S L A P A I K A G F G G A ttggaatggctacttccacgactgaaagagttaggagaatggttacagaaggcgagacggcgagaaggcgaaagagttcggtcagtct L E W L L P R L K E L G E W L Q K A G E K A K E F G Q S V G S K V S K L L E Q F G I S I G Q A G G S I G Q F I G aatgttctcgaaaggctaggaggcgcatttggaaaagtaggaggagtcatttcaattgctgtttcacttgtaacaaaattcggt N V L E R L G G A F G K V G G V I S I A V S L V T K F ctcgcatttctagggattacaggaccactcgggattgctattagtctgttagtttcatttttgacagcttgggctagaacaggt LAFLGITGPLGIAISLLVSFLTAWARTG gagttcaacgcagacggaattactcaagtattcgaaaacttgacaaacacaattcagtcgacggctgatttcatctctcaatac E F N A D G I T Q V F E N L T N T I Q S T A D F I S Q Y cttccagtctttgtcgaaaaaggaactcaaattttagttaagattattgaaggaattgcatctgctgttcctcaagtagttgaa L P V F V E K G T Q I L V K I I E G I A S A V P Q V V E gtgatttcacaagtcattgaaaatattgtgatgacaatttcgacagttatgcctcaattagtcgaagcaggaattaagatactt I S Q V I B N I V M T I S T V M P Q L V E A G I K I L gaagegettataaatggtettgtteaatetetteetaetateatteaageagetgtteaaattateaetgefftatteaatggt E A L I N G L V Q S L P T I I Q A A V Q I I T Ā L F N G  $\tt cttgttcaggcacttcctacgcttattcaagcaggtcttcaaattttgtcagctctcataaacggactagttcaagcgcttccg$ LVQALPTLIQAGLQILSALINGLVQALP gcaattattcaagcagctgttcaaattatcatgtcgcttgttcaagcactaattgaaaacttgcctatgataatcgaagcagcg A I I Q A A V Q I I M S L V Q A L I E N L P M I I E A A

atgcagattataatgggtctagtcaacgcactgattgaaaatataggacctatcttagaagcagggattcaaattctaatggct34654 M O I I M G L V N A L I E N I G P I L E A G I Q I L M A 757 ttaatcgagggacttattcaagtgcttcctgaactaattacagcagcgattcaaatcattacttcactattagaagcaatcttg 34738 LIEGLIQV LPELITAAIQIITS LLEAIL 785 tegaacetteeteaacttetagaageeggagttaaattgettttateacttetteaagggttgetaaatatgetteeteaacta
SNLPQLLBAGVKLLLSLLQGLLNMLPQL 34822 813 attgcaggggctttgcaaatcatgatggcacttcttaaagcagttatcgacttcgtccctaaacttcttcaagcaggtgttcaa I A G A L Q I M M A L L K A V I D F V P K L L Q A G V Q 34906 841 cttcttaaggcattgattcaaggtattgcttcacttctcggctcacttttatcgacagctggaaacatgctttcatcattagtt L L K A L I Q G I A S L L G S L L S T A G N M L S S L V 34990 869 35074 agcaagattgctagctttgtgggacagatggtttcaggaggtgcgaacctgattcgaaacctcattagtggtattgggtcaatggcgaacctgattcgaaacctcattagtggtattgggtcaatggcgaacctgattcgaaacctcattagtggtattgggtcaatggcgaacctgattcgaaacctgattcgaaacctcattagtggtattgggtattgggtattgggaacctgattcgaaacctcgaacctgattcgaaacctcattagtggtattgggtattgggtattgggaacctgaattcgaaacctcattagtggtattgggtattgggtattgggaacctgaattcgaaacctcgaattcgaaacctcattagtgggtattgggaacctgaattcgaaacctcgaattcgaaacctcattagtgggtattgggaacctgaattcgaaacctcgaattcgaaacctcaattagtggtattcgaattgggtattcaattggaagattggaaacctgaattcgaaacctcgaattcgaaacctcaattagtgggtattcgaaacctgaattcgaaacctcgaattcgaattcgaaacctcgaattcgaattcgaaacctcgaattcgaattcgaaacctcgaattcgaattcgaaacctcgaattcgaattcgaaacctcgaattcgaaacctcgaatS K I A S F V G Q M V S G G A N L I R N F I S G I G S M 897 attggttcagctgtctctaaaattggcagcatgggaacttcaattgtttctaaggttactggattcgctggacaaatggtaagc
I G S A V S K I G S M G T S I V S K V T G F A G Q M V S 35158 925 gcaggggtcaaccttgttcgaggatttatcaatggtatcagttccatggtaagttctgcggtaagtgcggcggctaatatggct 35242 A G V N L V R G F I N G I S S M V S S A V S A A A N M A 953 agcagtgcattaaatgccgttaagggattcttaggtattcactctccttcacgtgtcatggagcagatgggtatctatacgggt S S A L N A V K G F L G I H S P S R V M E Q M G I Y T G 35326 981 caagggttcgtaaatggtattggtaacatgattcgaactacacgtgacaaggctaaagaaatggctgaaactgttactgaagct 35410 Q G F V N G I G N M I R T T R D K A K E M A E T V T E A 1009 ctcagcgacgtgaagatggatattcaagaaaatggagttatagaaaaggttaaatcagtttacgaaaagatggctgaccaactt L S D V K M D I Q E N G V I B K V K S V Y E K M A D Q L 35494 1037 cctgaaactcttccagctcctgatttcgaagatgttcgtaaagcagccggttcgcctcgagtggacttgttcaatacaggaagt 35578 PETLPAPD FEDVRKAAGSPRVDL FNTGS 1065 gacaaccctaaccaacctcagtcacaatctaaaaacaatcaaggcgagcaaaccgttgtcaacattggaacaatcgtagttcga 35662 D N P N Q P Q S Q S K N N Q G E Q T V V N I G T I V V R 1093 35746 aacaatgacgacgttgacaaactgtcgagaggattgtataatagaagtaaagtaactctatcagggtttggtaacattgtaaca 1121 N N D D V D K L S R G L Y N R S K E T L S G F G N I ccgtaa 35835 35830

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2012
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2096
          K L L V E S *
29
dp10RF250
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23837
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23921
          KSNHAL
29
dp10RF251
          {\tt atggaaataattagtcttaccgtctgcgcctggcttcccgggtatcccttgagctccgtcattccccttccatttcgtccatgt}
39205
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          ataggetgeagggtettttga 39101
39121
          IGCRVF
29
 dp10RF252
          gtgttgtataggtcgaaactaattttgcatattttctatatttcaaaagtgcttttgagatatcgttateaaaatgctcgacaa
54771
          V L Y R S K L I L H I F Y I S K V L L R Y R Y Q N A R Q
1
          tactttcgcctgttcctctag 54667
54687
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 dp10RF253
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 56255
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dp10RF254
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48479
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48395
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       PDSMPK *
dp10RF255
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9572
9488
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15289
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15373
       H L K K F
29
dp10RF257
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28216
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28300
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44023
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44107
       gcgatggttcaatggtaa 44124
           V Q W
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4298
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4382
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       24746
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24830
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dp10RF261
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372
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       LNLNH *
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       M K I L A S S S F E V F E I I S F T C L I V G S S R P F
       aacaagtcttctaattga 26951
26968
       NKSSN
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dp10RF264
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6139
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6055
       tgtgaaacgtcttcataa 6038
29
       CETSS *
dp10RF265
4801
       \tt gtgaataaagtcaagegtttttgtataaaaagttcattttttttaaaaaaaataagagegaaaagetcttatctaaaatagte
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       gacgttgacgatttttaa 4700
4717
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       DVDDF
dp10RF266
50220
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       M P.V L P S S C K H F I N S P R L T L S R S S H Y D N Q atcctcaccaggaagtaa 50119
1
                                                               50136
       ILTRK
29
dplORF267
47367
       \verb|atggtcaaggtctgttctaggttcaggaagaacaaacgggaagtgaatgttattttcttcagcgaagtcttttgcttcatacca|\\
       M V K V C S R F R K N K R E V N V I F F S E V F C F I P
       aacattaatcgtagatag 47266
47283
dp10RF268
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12621
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                                        M S I S V L C L T M D S T T D A S T F F N R D S L S N S
12537
                                        ttgtcaattctagagtaa 12520
29
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 dp10RF269
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1
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29
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50792
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                                        M R L L C F I F V T V L T D F L L A N L P T R I H T S K
19655
                                       gctttttgtcagccttag 19638
29
 dp10RF272
                                       gtggtcaagtctgtcaatgaatgtacctgcgattttcttgacgtgataaaagtcaacaaccatcccttgactcgaaccgtggtc V V K S V N B C T C D F L D V I K V N N H P L T R T V V ataagttccgcctgctaa 1455
1556
1
1472
29
                                       ISSAC .
dp10RF273
56256
                                       {\tt atggatttcattaggactgagtcctcttggaattggaacggttgcatatatagatattccgtcagccgtactaggccaagttct}
                                       M D F I R T E S S W N W N G C I Y R Y S V S R T R P S S
                                      56340
29
56424
```

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#### Table 31

Query= sid|114822|lan|dp1ORF001 Phage dp1 ORF|36698-40390|2 (1230 letters) >gi|928828 (L44593) ORF1904; putative [Lactococcus lactis phage BK5-T] Length = 1904 Score = 427 bits (1086), Expect = e-118 Identities = 226/475 (47%), Positives = 281/475 (58%), Gaps = 45/475 (9%) Query: 395 AESGKYIGVLNTNKKPSELVPDDFTWIRLEGPKGDAGLPGAPGRDGVDGVPGKSGVGIAD 454
A+ YIG + P D+TW + +G+ G GA G+DGV GK GVGI
Sbjct: 820 ADYPSYIGQYTDFIQYDSAKPSDYTWSLI---RGNDGKDGATGKDGV---AGKDGVGIKT 873 Query: 455 TAITYAVSVSGTQEPENGWSEQVPELIKGRFLWTKTFWRYTDGSHETGYSVAYIGQDGNS 514 T ITYA+S SGT +P GW+ QVP L+KG++LWTKT W YTD S ETGYSV YI +DGN+ Sbjct: 874 TVITYALSSSGTDKPNTGWTSQVPTLVKGQYLWTKTVWTYTDSSSETGYSVTYIAKDGNN 933 Query: 515 GKDGIAGKDGVGIAATEVMYASSPSATEAPAGGWSTQVPTVPGGQYLWTRTRWRYTDQTD 574 G DGIAGKDGVGI T + YA S T APA GW++QVP VP GQ+LWT+T W YTD T Sbjct: 934 GNDGIAGKDGVGIKKTTITYAVGTSGTTAPASGWNSQVPNVPAGQFLWTKTVWTYTDNTS 993 Query: 575 EIGYSVSRMGEQGPKGDAGR---DGIAGKNGIGLKSTSVSYGISPTDSAIP-GVWASQVP 630 E GYSV+ MG +G KGD G +GIAGK+G G+K+T+++Y SP + P G W++ VP Sbjct: 994 ETGYSVAMMGVKGDKGDFGNNGTNGIAGKDGKGIKATAITYQASPNGTTAPTGTWSASVP 1053 Query: 631 SLIKGQYLWTRTIWTYTDSTTETGYQKTYIPKDGNDGKNGIAGKDGVGIKSTTITYAGST 690 + KG +LWTRTIWTYTD+TTETGY Y+ +GN+G +G GKDG GIK+TTITYAGST
Sbjct: 1054 PVAKGSPLWTRTIWTYTDNTTETGYAVAYMGTNGNNGHDGPPGKDGTGIKTTTITYAGST 1113 SGT P + WTS +P V G +LWTKTVW YTD+TSETGYSV+ +G Sbjct: 1114 SGTTPPNNGWTSTVPTVAEGNYLWTKTVWTYTDNTSETGYSVAMMG-----VKGDKGDP 1167 Query: 751 XXXXXXXXXADGRS-QYTHLAFSNSPNGEGFSHTDSGRAYVGQYQDFNPVHSKDPAAYT 809 DG+ + T + + SPNG A G + Sbjct: 1168 GNNGTNGIAGKDGKGIKATAITYQASPNGT-----TAPTGTWSASVPPVAKGSFLWT 1219 Query: 810 WTKW------KGNDGAQGIPGKPGADGKTNYFHIAYASSADGS 846 GN+G G PGK G KT I YA S G+ Sbjct: 1220 RTIWTYTDNTTETGYAVAYMGTNGNNGHDGPPGKDGTGIKTT--TITYAGSTSGT 1272 Score = 396 bits (1007), Expect = e-109 Identities = 208/449 (46%), Positives = 260/449 (57%), Gaps = 42/449 (9%) Query: 421 IRLEGPKGDAGLPGAPGRDGVDGVPGKSGVGIADTAITYAVSVSGTQEPENGWSEQVPEL 480
+ + G KGD G PG +G +G+ GK G GI TAITY S +GT P WS VP + Sbjct: 1155 VAMMGVKGDKG---DPGNNGTNGIAGKDGKGIKATAITYQASPNGTTAPTGTWSASVPPV 1211 Query: 481 IKGRFLWTKTFWRYTDGSHETGYSVAYIGQDGNSGKDGIAGKDGVGIAATEVMYASSPSA 540 KG FLWT+T W YTD + ETGY+VAY+G +GN+G DG GKDG GI T + YA S S Sbjct: 1212 AKGSFLWTRTIWTYTDNTTETGYAVAYMGTNGNNGHDGFPGKDGTGIKTTTITYAGSTSG 1271 Query: 541 TEAPAGGWSTQVPTVPGGQYLWTRTRWRYTDQTDEIGYSVSRMGEQGPKGDAGR---DGI 597 T P GW++ VPTV G YLWT+T W YTD T E GYSV+ MG +G KGD G Sbjct: 1272 TTPPNNGWTSTVPTVAEGNYLWTKTVWTYTDNTSETGYSVAMMGVKGDKGDPGNNGTNGI 1331 Query: 598 AGKNGIGLKSTSVSYGISPTDSAIP-GVWASQVPSLIKGQYLWTRTIWTYTDSTTETGYQ 656 AGK+G G+K+T+++Y SP + P G W++ VP + KG +LWTRTIWTYTD+TTETGY Sbjct: 1332 AGKDGKGIKATAITYQASPNGTTAPTGTWSASVPPVAKGSFLWTRTIWTYTDNTTETGYA 1391 Query: 657 KTYIPKDGNDGKNGIAGKDGVGIKSTTITYAGSTSGTVAPTSNWTSAIPNVQPGFFLWTK 716 Y+ +GN+G +G GKDG GIK+TTITYAGSTSGT P + WTS +P V G +LWTK Sbjct: 1392 VAYMGTNGNNGHDGFPGKDGTGIKTTTITYAGSTSGTTPPNNGWTSTVPTVAEGNYLWTK 1451 Query: 717 TVWNYTDDTSETGYSVSKIGETXXXXXXXXXXXXXXXXXXXXXXXXADGRS-QYTHLAFSNS 775 TVW YTD+TSETGYSV+ +G DG+ + T + + S

Sbjct: 1452 TVWTYTDNTSETGYSVAMMG------VKGDKGDPGNNGTNGIAGKDGKGIKATAITYOAS 1505

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Query: 776 PNGEGFSHTDSGRAYVGQYQDFNPVHSKDPAAYTWTKW------KGND 817
             PNG
                           A G + P +K
                                                +T T W
                                                                             GN+
Sbjct: 1506 PNGT-----TAPTGTWSASVPPVAKGSFLWTRTIWTYTDNTTETGYAVAYMGTNGNN 1557
Query: 818 GAQGIPGKPGADGKTNYFHIAYASSADGS 846
G G PGK G KT I YA S G+
Sbjct: 1558 GHDGFPGKDGTGIKTT--TITYAGSTSGT 1584
 Score = 384 bits (977), Expect = e-105
Identities = 179/322 (55%), Positives = 222/322 (68%), Gaps = 7/322 (2%)
Query: 421 IRLEGPKGDAGLPGAPGRDGVDGVPGKSGVGIADTAITYAVSVSGTQEPENGWSEQVPEL 480
+ + G KGD G PG +G +G+ GK G GI TAITY S +GT P WS VP +
Sbjet: 1311 VAMMGVKGDKG---DPGNNGTNGIAGKDGKGIKATAITYQASPNGTTAPTGTWSASVPPV 1367
Query: 481 IKGRFLWTKTFWRYTDGSHETGYSVAYIGQDGNSGKDGIAGKDGVGIAATEVMYASSPSA 540
              KG FLWT+T W YTD + ETGY+VAY+G +GN+G DG GKDG GI T + YA S S
Sbjct: 1368 AKGSFLWTRTIWTYTDNTTETGYAVAYMGTNGNNGHDGFPGKDGTGIKTTTITYAGSTSG 1427
Query: 541 TEAPAGGWSTQVPTVPGGQYLWTRTRWRYTDQTDEIGYSVSRMGEQGPKGDAGR---DGI 597
T P GW++ VPTV G YLWT+T W YTD T E GYSV+ MG +G KGD G +GI
Sbjct: 1428 TTPPNNGWTSTVPTVAEGNYLWTKTVWTYTDNTSETGYSVAMMGVKGDKGDPGNNGTNGI 1487
Query: 598 AGKNGIGLKSTSVSYGISPTDSAIP-GVWASQVPSLIKGQYLWTRTIWTYTDSTTETGYQ 656
AGK+G G+K+T+++Y SP + P G W++ VP + KG +LWTRTIWTYTD+TTETGY
Sbjct: 1488 AGKDGKGIKATAITYQASPNGTTAPTGTWSASVPPVAKGSFLWTRTIWTYTDNTTETGYA 1547
Query: 657 KTYIPKDGNDGKNGIAGKDGVGIKSTTITYAGSTSGTVAPTSNWTSAIPNVQPGFFLWTK 716
               Y+ +GN+G +G GKDG GIK+TTITYAGSTSGT P + WTS +P V G +LWTK
sbjct: 1548 VAYMGTNGNNGHDGFPGKDGTGIKTTTITYAGSTSGTTPPNNGWTSTVPTVAEGNYLWTK 1607
Query: 717 TVWNYTDDTSETGYSVSKIGET 738
             TVW YTD++ ETGYSV K+G T
Sbjct: 1608 TVWAYTDNSFETGYSVGKMGNT 1629
 Score = 201 bits (507), Expect = 2e-50
 Identities = 121/297 (40%), Positives = 156/297 (51%), Gaps = 19/297 (6%)
Query: 421 IRLEGPKGDAGLPGAPGRDGVDGVPGKSGVGIADTAITYAVSVSGTQEPENGWSEQVPEL 480
+ + G KGD G PG +G +G+ GK G GI TAITY S +GT P WS VP +
Sbjct: 1467 VAMMGVKGDKG---DPGNNGTNGIAGKDGKGIKATAITYQASPNGTTAPTGTWSASVPPV 1523
Query: 481 IKGRFLWTKTFWRYTDGSHETGYSVAYIGQDGNSGKDGIAGKDGVGIAATEVMYASSPSA 540
              KG PLWT+T W YTD + ETGY+VAY+G +GN+G DG GKDG GI T + YA S S
Sbjct: 1524 AKGSFLWTRTIWTYTDNTTETGYAVAYMGTNGNNGHDGFPGKDGTGIKTTTITYAGSTSG 1583
Query: 541 TEAPAGGWSTQVPTVPGGQYLWTRTRWRYTDQTDEIGYSVSRMGEQGPKGDAGRDGIAGK 600
             T P GW++ VPTV G YLWT+T W YTD + E GYSV +MG GP AG +G GK
Sbjct: 1584 TTPPNNGWTSTVPTVAEGNYLWTKTVWAYTDNSFETGYSVGKMGNTGP---AGSNGNPGK 1640
Query: 601 NGIGLKSTSVSYGISPTDSAIPGVWASQVPSLIKG-QYLWTRTIWTYTDSTTE--TGYQK 657
                  + T+ G++ S + + ++ G +Y W W +
Sbjct: 1641 VVSDTEPTTKFKGLTWKYSGVVDMPLGNGTKILAGTEYYWNGNNWALYEINAHNINGDNL 1700
Query: 658 TYIPKDGNDGK-NGIAGKDGVGIKSTTITYAGS----TSGTVAPTSNWTSAIPNVQ 708
+ DGK I G +GV + T T GS +S + T N T AI N Q
Sbjct: 1701 SVTNGTFKDGKIESIWGSNGV---NGTTTIEGSHLQIHSSDSTTNTEN-TLAIDNRQ 1753
Query= sid | 114823 | lan | dp1ORF002 Phage dp1 ORF | 32386-35835 | 1
          (1149 letters)
>dbj|BAA31888| (AB009866) orf 15 (bacteriophage phi PVL)
 Score = 280 bits (709), Expect = 3e-74
 Identities = 157/465 (33%), Positives = 257/465 (54%), Gaps = 28/465 (6%)
Query: 40 QIGSALTGLGKGLTTAVTLPLMGFAAASIKVGNEFQAQMSRVQAIAGATAEELGRMKTQA 99
            +IG+++ +G+ +T VT P++ A + K G EF M +V+A +GAT EE +K +A
Sbjct: 151 EIGNSMKNVGRNMTMYVTAPVVAGFAVAAKKGIEFDDSMRKVKATSGATGEEFRALKKKA 210
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WO 00/32825 PCT/IB99/02040

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++GA T FSA ++A+ + +A AG+ ++M+ + GV+DL
Sbjct: 211 REMGATTKFSASDSAEALNYMALAGWDSKOMMEGLSGVMDLAAASGEELGAVSDIVTDGL 270
Query: 160 RAFGLEANQAGHVADVFARAAADTNAETSDMAEAMKYVAPVAHSMGLSLEETAASIGIMA 219
AFGL+A +GH+ADV A+ ++ N + + EA KYVAPVA ++G ++E+T+ +IG+M+
Sbjct: 271 TAFGLKAKDSGHLADVLAQTSSKANTDVRGLGEAFKYVAPVAGALGYTIEDTSIAIGLMS 330
Query: 220 DAGIKGSQAGTTLRGALSRIAKPTKAMVKSMQELGVSFYDANGNMIPLREQIAQLKTATA 279
            +AGIKG +AGT LR + ++ PT+AM M+ LG+S D+NG MIP+R+ + QL+
Sbjct: 331 NAGIKGEKAGTALRTMFTNLSSPTRAMGNEMERLGISITDSNGKMIPMRKLLDQLREKFK 390
Query: 280 GLTQEERNRHLVTLYGQNSLSGMLALLDAGPEKLDKMTNALVNSDGAAKEMAETMQDNLA 339
                      T++G+ ++SG LA+++A E K+T ++ +S GA+K MA+TM+ L
Sbjct: 391 HLSKDQQASSAATIFGKEAMSGALAIINASDEDYQKLTKSIDSSTGASKRMADTMESGLG 450
Query: 340 SKIEQMGGAFESVAIIVQQILEPALAKIVGAITKVLEAFVNMSPIGQKMVVIFAGMVAAL 399
K+ + E +A+ + +EPAL IV A +KV+ + Q VV F VA L
Sbjct: 451 GKLRTLRSQLEELALTIYDRIEPALKIIVSAPSKVVTWVTKLPTSIQLAVVGFGLFVAVL 510
Query: 400 GPLLLIAGM------ VMTTIVKLRIAIQFLGPAFMGTMGTIAGVIAIF----- 441
GPL+ + G+ MT + L I + F IA ++ +F
Sbjct: 511 GPLVFMFGLFISVMGNAMTVLGPLLINVNKASGLFAFLRTKIASLVKLFPILGVSISSLT 570
Query: 442 ------YALVAV---FMIAYTKSERFRNFINSLAPAIKAGFGGA 476
ALV + F AY +SE FRN +N + F A
Sbjct: 571 LPITLIVGALVGIGIAFYQAYKRSETFRNIVNQAISGVANAFKAA 615
Query= sid|114824|lan|dp10RF003 Phage dp1 ORF|53538-55877|3
          (779 letters)
>ap|P43741|DP01_HAEIN DNA POLYMERASE I (POL I) >gi|1074025|pir||E64098 DNA polymerase I
            (polA) homolog - Haemophilus influenzae (strain Rd KW20)
            >gi|1573871 (U32767) DNA polymerase I (polA)
            (Haemophilus influenzae Rd)
            Length = 930
 Score = 191 bits (481), Expect = 1e-47
 Identities = 148/553 (26%), Positives = 262/553 (46%), Gaps = 60/553 (10%)
Query: 63 RLELITERAKLEQYVDKMIEDGIGSIDVETDGLDTIHDELAGVCLYSPSQKGIYAPVNHV 122
            + E + +A L ++++K+ + ++D ETD LD + L G+
Sbjct: 333 KYETLLTQADLTRWIEKLNAAKLIAVDTETDSLDYMSANLVGISFALENGEAAYLPLQLD 392
Query: 123 SNMTKMRIKNQISPEFMKKMLQRIVDSGIPVIYHNSKFDMKSIYWRLGVKMNEPAWDTYL 182
++ + +K +L+ + I I N KFD +SI+ R G+++ +DT L
Sbjct: 393 YLDAPKTLEKSTALAAIKPILE---NPNIHKIGQNIKFD-ESIFARHGIELQGVEFDTML 448
Query: 183 AAMLLNENESHSLKSLHSKYVRNEENAEVAKFNDLFKGIPFSLIPPDVAYMYAAYDPLQT 242
+ IN H++ L +Y+ +E A + + F+ IP + A YAA D T
Sbjct: 449 LSYTLNSTGRHNMDDLAKRYLGHETIAFESLAGKGKSQLTFNQIPLEQATEYAAEDADVT 508
Query: 243 FELYEFQEQYLTPGTEQCEEYNLEKVSWVLHNIEMPLIKVLFDMEVYGVDLDQDKLAEIR 302
                                              +E+PL+ VL ME GV +D D L
                              EΥ
Sbjct: 509 MKLQQALWLKLQEEPTLVELYK------TMELPLLHVLSRMERTGVLIDSDALFMQS 559
Query: 303 EQFTANMNEAEQEFQQLVSEWQPEIEELRQTNFQSYQKLEMDARGRVTVSISSPTQLAIL 362
Query: 363 FYDIMGLKSPERDKPRG---TGESIVEH--FDNDISXXXXXXXXXXXXXXXXXXTTTT-LDQHL 416 +D + L ++ P+G T E ++E + +++ STYT L Q +
Sbjct: 593 LFDKLELPVLQKT-PKGAPSTNEEVLEELSYSHELPKILVKHRGLSKLKSTYTDKLPQMV 651
Query: 417 AKPDNRIHTTFKQYGAKTGRMSSENPNLQNIPSRGE-GAVVRQIFAASEGHYIIGSDYSQ 475
R+HT++ Q TGR+SS +PNLQNIP R E G +RQ F A EG+ I+ +DYSQ Sbjct: 652 NSQTGRVHTSYHQAVTATGRLSSSDPNLQNIPIRNEEGRHIRQAFIAREGYSIVAADYSQ 711
Query: 476 QEPRSLAELSGDESMRHAYEQNLDLYSVIGSKLYGVPYEECLEFYPDGTTNKEGKLRRNS 535
E R +A LSGD+ + +A+ Q D++ ++++GV +E T+++ R +
Sbjct: 712 IELRIMAHLSGDQGLINAFSQGKDIHRSTAAEIFGVSLDE------VTSEQ----RRN 759
 Query: 536 VKSVLIGLMYGRGANSIAEQMNVSVKEANKVIEDFFTEFPKVADYIIFVQQQAQDLGYVQ 595
             K++ GL+YG A ++ Q+ +S +A K ++ +F +P V ++ ++++A+ GYV+
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Sbjct: 760 AKAINFGLIYGMSAFGLSRQLGISRADAQKYMDLYFQRYPSVQQFMTDIREKAKAQGYVE 819
Ouerv: 596 TATGRRRRLPDMS 608
           T GRR LPD++
Sbjct: 820 TLFGRRLYLPDIN 832
 Score = 46.9 bits (109), Expect = 5e-04
 Identities = 34/123 (27%), Positives = 66/123 (53%), Gaps = 16/123 (13%)
Query: 663 EIKDQAKAEGI------LIKDNGGKIADAQRQCLNSVIQGTAADMTKYAMIKV 709
           +I+++AKA+G
                                  + N + A+R +N+ +QGTAAD+ K AMIK+
Sbjct: 807 DIREKAKAQGYVETLFGRRLYLPDINSSNAMRRKGAERVAINAPMQGTAADIIKRAMIKL 866
Query: 710 HNDAELKELGFHLMIPVHDELLGEVPIKNAKRGAERLTEVMIEAAKDIISLPMKCDPSIV 769
++ + +++ VHDEL+ EV + E++ M EAA +++ +P+ + +
Sbjct: 867 -DEVIRHDPDIEMIMQVHDELVFEVRSEKVAFFREQIKQHM-EAAAELV-VPLIVEVGVG 923
Query: 770 ERW 772
Sbjct: 924 QNW 926
Query= sid|114825|lan|dp1ORF004 Phage dp1 ORF|40401-42440|3
         (679 letters)
>emb|CAB07981| (Z93946) hypothetical protein [bacteriophage Dp-1]
           Length = 532
 Score = 1011 bits (2585), Expect = 0.0
 Identities = 497/499 (99%), Positives = 498/499 (99%)
          MTKFINSYGPLHLNLYVEQVSQDVTNNSSRVSWRATVDRDGAYRTWTYGNISNLSVWLNG 60
           MTKFINSYGPLHLNLYVEQVSQDVTNNSSRVSWRATVDRDGAYRTWTYGNISNLSVWLNG
Sbict: 1
          MTKFINSYGPLHLNLYVEQVSQDVTNNSSRVSWRATVDRDGAYRTWTYGNISNLSVWLNG 60
Query: 61 SSVHSSHPDYDTSGEEVTLASGEVTVPHNSDGTKTMSVWASFDPNNGVHGNITISTNYTL 120
           SSVHSSHPDYDTSGEEVTLASGEVTVPHNSDGTKTMSVWASFDPNNGVHGNITISTNYTL
Sbjct: 61 SSVHSSHPDYDTSGEEVTLASGEVTVPHNSDGTKTMSVWASFDPNNGVHGNITISTNYTL 120
Query: 121 DSIPRSTQISSFEGNRNLGSLHTVIFNRKVNSFTHQVWYRVFGSDWIDLGKNHTTSVSFT 180
           DSIPRSTQISSFEGNRNLGSLHTVIFNRKVNSFTHQVWYRVFGSDWIDLGKNHTTSVSFT
Sbjct: 121 DSIPRSTQISSFEGNRNLGSLHTVIFNRKVNSFTHQVWYRVFGSDWIDLGKNHTTSVSFT 180
Query: 181 PSLDLARYLPKSSSGTMDICIRTYNGTTQIGSDVYSNGWRFNIPDSVRPTFSGISLVDTT 240
           PSLDLARYLPKSSSGTMDICIRTYNGTTQIGSDVYSNGWRFNIPDSVRPTFSGISLVDTT
Sbjct: 181 PSLDLARYLPKSSSGTMDICIRTYNGTTQIGSDVYSNGWRFNIPDSVRPTFSGISLVDTT 240
Query: 241 SAVRQILTGNNFLQIMSNIQVNFNNASGAYGSTIQAFHAELVGKNQAINENGGKLGMMNF 300
           SAVRQILTGNNFLQIMSNIQVNFNNASGAYGSTIQAFHAELVGKNQAINENGGKLGMMNF
Sbjct: 241 SAVRQILTGNNFLQIMSNIQVNFNNASGAYGSTIQAFHAELVGKNQAINENGGKLGMMNF 300
Query: 301 NGSATVRAWVTDTRGKQSNVQDVSINVIEYYGPSINFSVQRTRQNPAIIQALRNAKVAPI 360
           NGSATVRAWVTDTRGKQSNVQDVSINVIEYYGPSINFSVQRTRQNPAIIQALRNAKVAPI
Sbjct: 301 NGSATVRAWVTDTRGKQSNVQDVSINVIEYYGPSINFSVQRTRQNPAIIQALRNAKVAPI 360
Query: 361 TVGGQQKNIMQITFSVAPLNTTNFTEDRGSASGTFTTISLMTNSSANLAGNYGPDKSYIV 420
           TVGGQQKNIMQITFSVAPLNTTNFTEDRGSASGTFTTISL+TNSSANLAGNYGPDKSYIV
Sbjct: 361 TVGGQQKNIMQITFSVAPLNTTNFTEDRGSASGTFTTISLLTNSSANLAGNYGPDKSYIV 420
Query: 421 KAKIQDRFTSTEFSATVATESVVLNYDKDGRLGVGKVVEQGKAGSIDAAGDIYAGGRQVQ 480
           KAKIQDRFTSTEFSATV TESVVLNYDKDGRLGVGKVVEQGKAGSIDAAGDIYAGGRQVQ
Sbjct: 421 KAKIQDRFTSTEFSATVPTESVVLNYDKDGRLGVGKVVEQGKAGSIDAAGDIYAGGRQVQ 480
Query: 481 QFQLTDNNGALNRGQYNDV 499
           QFQLTDNNGALNRGQYNDV
Sbjct: 481 QFQLTDNNGALNRGQYNDV 499
Query= sid|114827|lan|dp10RF006 Phage dp1 0RF|45296-46987|2
                                                                                 ______
         (563 letters)
>gb|AAD18987| (AE001666) SWI/SNF family helicase_2 [Chlamydia pneumoniae]
           Length = 1166
 Score = 171 bits (429), Expect = 1e-41
 Identities = 150/522 (28%), Positives = 254/522 (47%), Gaps = 55/522 (10%)
```

```
SSNNFE-LPYKYFNNVIDALDEWELHIFGELDKDVQDYIDSRNRIASSSNEQFSFKTTPF 104
              S + FE LP + ++ + L E + I GE++ D QD
Sbjct: 659 SLDQFEALPVNF--SMSERLIEIQKQIRGEIEFDFQD------VPQQIQATLRSYQTEG 709
Query: 105 AHQVECFEYAQEHPCFLLGDEQGLGKTKQAIDIAVSRKASFKH--CLIVCCISGLKWNWA 162
                         + H +L D+ GLGKT QAI IAV++
                                                             K C ++ C + L +NW
Sbjct: 710 VHWLE--RLRKMHLNGILADDMGLGKTLQAI-IAVTQSKLEKGSGCSLIVCPTSLVYNWK 766
Query: 163 KEVGIHSNESAHILGSRVTKDGKLVIDGV-SKRAEDLLGGHDEFFLITNIETLRDAVFIK 221
                                        LVIDGV S+R + L D IT+ L+ V
               +E
                    + E
Sbict: 767 EEFRKFNPEFR------TLVIDGVPSORRKOLTALADRDVAITSYNLLOKDV--- 812
Query: 222 YLNELTKSGEIGMVIIDEIHKCKNPSSKQGASIQKLQSYYKMGLTGTPLMNNPIDVFNVM 281
                            V++DE H KN +++ S++ +QS +++ LTGTP+ N+ +++++
                  EL KS
              ---ELYKSFRFDYVVLDEAHHIKNRTTRNAKSVKMIQSDHRLILTGTPIENSLEELWSLF 869
Sbjct: 813
Query: 282 KWLGAEHHTLTQFKERYCIVDQFNQITGYR----NLAELRELVNDYMLRRTKEEVL-DL 335
                      L +R+ V ++ + Y N+ L++ V+ ++LRR KE+VL DL
              DFLMPG---LLSSYDRF--VGKYIRTGNYMGNKADNMVALKKKVSPFILRRMKEDVLKDL 924
Query: 336 PEKIRVTEYVDMNSKQSKIY------KEVLTKLVQEIDKVKLMPNPLAETIRLRQATGN 388
P + + + Q ++Y K+ L++LV++ ++ + LA RL+Q +
Sbjct: 925 PPVSEILYHCHLTESQKELYQSYAASAKQELSRLVKQEGFERIHIHVLATLTRLKQICCH 984
Query: 389 PSILTTQDVK---SCKFERCIEIVEECIQQGKSCVIFSNWEKVIEPLAKIL-SKTVKCNL 444
P+I + S K++ ++++ + G V+FS + K++ + K L S+ +
Sbjct: 985 PAIFAKDAPBPGDSAKYDMLMDLLSSLVDSGHKTVVFSQYTKMLGIIKKDLESRGIPFVY 1044
Query: 445 VTGETADKFNEIEEFMNHRKASVILGTIGALGTGFTLTKADTVIFLDSPWTRAEKDQAED 504
+ G T ++ + + +F V L ++ A GTG L ADTVI D W A ++QA D
Sbjct: 1045 LDGSTKNRLDLVNQFNEDPSLLVFLISLKAGGTGLNLVGADTVIHYDMWWNPAVENQATD 1104
Query: 505 RCHRIGAKSSVTIYTLVAKGTVDERIEDLIERKGELADYIVD 546
              R HRIG SV+ Y LV T++E+I L RK L
Sbjct: 1105 RVHRIGQSRSVSSYKLVTLNTIEEKILTLQNRKKSLVKKVIN 1146
Query= sid | 114828 | lan | dp10RF007 Phage dp1 ORF | 22230-23621 | 3
           (463 letters)
>gi|2444105 (U88974) ORF26 [Streptococcus thermophilus temperate bacteriophage
             012051
             Length = 411
 Score = 88.9 bits (217), Expect = 7e-17 Identities = 80/315 (25%), Positives = 133/315 (41%), Gaps = 48/315 (15%)
Query: 139 QGVTLAGIFCDEVALMPESFVNQATGRCSVTGSKMWFSCNPANPNHYFKKNWIDKQVEKR 198
+G T G + +E +L E + RCS G+++ + NP NPNH+ +++I K + +
Sbjct: 121 RGFTAFGAYVNEASLANELVFKEIISRCSGDGARVVWDSNPDNPHWLNRDYIGKN-DGK 179
Query: 199 ILYLHFTMDDNPSLT----DSIKRRYEKMYAGVFRKRFILGLWVTADGLVYSMFNEEQHV 254
I+ F +DDN L+ DSIK K G F R ILGLW A+G +Y+ ++ +HV
Sbjct: 180 IIDFSFKLDDNTFLSKRYIDSIKAATPK---GKFYDRDILGLWTVAEGAIYADYDSKIHV 236
Query: 255 KKLNIEFDRLFVAGDFGIYNATTFGLYGFSKRHKRYHLIESYYHSGREAEEQLTEADVNS 314
E R F D+G + + + G ++L++ +B + + +A
Sbjct: 237 VDELPEMKRYFGGIDWGYTHYGSIVIVG-EGVDNNFYLVDGVAAQFKEIDWWVEQA---- 291
Query: 315 NIQFSSVLQKTTKEYANDLVDMIRGKQIEYIILDPSASAMIVELQKHPYIAR---KNIPI 371
                                                                 + ++AR +
                      +KT YN
Sbjct: 292 -----RKLTGIYGN------IPFYADSARPEHVARFENEGFDI 323
Query: 372 IPARNDVTLGISFHAELLAENRFTLDPSNT-HDIDEYYAYSWDSKASQTGEDRVIKEHDH 430
                   V GI A+L E + +
                                                   DE Y Y W ++ +D +KE D
Sbjct: 324 MNANKSVIAGIELIAKLFKEKKLYVKRGFVPRFFDEIYQYRWKENST---KDEPLKEFDD 380
Query: 431 CMDRNRYACLTDALI 445
                                                                                               +D RYA +D +I
Sbjct: 381 VLDSVRYAIYSDYVI 395
Query= sid | 114829 | lan | dp10RF008 Phage dp1 ORF | 49624-50961 | 1
           (445 letters)
```

>gb|AAD19901| (AF100420) DnaB replication fork helicase [Thermus aquaticus]

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Length = 444
```

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Score = 67.5 bits (162), Expect = 2e-10
 Identities = 69/248 (27%), Positives = 111/248 (43%), Gaps = 14/248 (5%)
Query: 147 GERLGISTGFEXXXXXXXXXXXXXXIVIMARPGQGKS-WTIDKMLATAWKNGHDVLLYS 205
GE G+ TGF+ I I ARP GK+ + + A K G V +YS
Sbjct: 178 GEVAGVRTGFKELDQLIGTLGPGSLNI-IAARPAMGKTAFALTIAQNAALKEGVGVGIYS 236
Query: 206 GEMSEMQVGARIDTILSNVSINSITKGIWNDHQFEKYEDHIQAMTRAENSLVVVTPFMIG 265
            EM Q+ R+ +++N+ G D F+ D
                                                      ++EA
Sbict: 237 LEMPAAOLTLRMMCSEARIDMNRVRLGOLTDRDFSRLVDVASRLSEAP-IYIDDTPDLTL 295
Query: 266 GKNLTPAILDSMISKYRPSVVGIDQLSLMS--ESYPSREQKRIQYANITMDLYKISAKYG 323
                      ++S+ + ++ ID L LMS S S E ++ + A I+ L ++ + G
Sbjct: 296 ME--VRARARRLVSQNQVGLIIIDYLQLMSGPGSGKSGENRQQEIAAISRGLKALARELG 353
Query: 324 IPIVLNVQAGRSAKTEGAESMELEHIAESDGVGQNASRVIAMKRD-----EKSGILEL 376
           IPI+ Q R+ + L + ES + Q+A V+ + RD
Sbjct: 354 IPIIALSQLSRAVEARPNKRPMLSDLRESGSIEQDADLVMFIYRDEYYNPHSEKAGIAEI 413
Query: 377 SVVKNRYG 384
            VKRG
Sbjct: 414 IVGKQRNG 421
Query= sid | 114831 | lan | dp1ORF010 Phage dp1 ORF | 8699-9859 | 2
         (386 letters)
>gi|2760912 (AF037258) RecA protein [Chlorobium tepidum]
           Length = 346
 Score = 133 bits (331), Expect = 2e-30
 Identities = 99/340 (29%), Positives = 164/340 (48%), Gaps = 66/340 (19%)
GGLPR RV E +GPESSGKTT AL + AQ
Sbjct: 67 GGLPRGRVTEIYGPESSGKTTLALHAIAEAQ-
Query: 104 AVKELEMQLDSLQEPLKIVYLDLENTLDTEWAKKIGVDVDNIWIVRPEMNSAEEILQYVL 163
+ L +D E+ D +A+K+GVD++ + + +PE S E+ L V
Sbjct: 101 GIAAL------VDAEHAFDPTYARKLGVDINALLVSQPE--SGEQALSIVE 143
Query: 164 DIFETGEVGLVVLDSLPYMVSQNLIDEELTKKAYAGISAPLTEFSRKVTPLLTRYNAIFL 223
             + +G V ++V+DS+ +V Q ++ E+
                                                + +++ RK+T +++ L
Sbjct: 144 TLVRSGAVDIIVIDSVAALVPQAELEGEMGDSVVGLQARLMSQALRKLTGAISKSSSVCL 203
Query: 224 GINQIREDMNSQYNA-YSTPGGKMWKHACAVRLKFRKGDYLDENGASLTRTARNPAGNVV 282
INQ+R+ + Y + +T GGK K +VKL RK + ++G L GN
Sbjct: 204 FINQLRDKIGVMYGSPETTTGGKALKFYSSVRLDIRKIAQI-KDGEELV------GNRT 255
Query: 283 ESFVEKTKAFKPDRKLVSYTLSYHDGIQIENDLVDVAVEFGVIQKAGAWFSIVDLETGEI 342
             V K K P K + + Y +GI + +L+D+AVEFG+I+K+GAWFS + G
Sbjct: 256 KVKVVKNKV-APPFKTAEFDILYGEGISVLGELIDLAVEFGIIKKSGAWFSYGTEKLG-- 312
Query: 343 MTDEDEEPLKFQGKANLVRRFKEDDYLFDMVMTAVHEIIT 382
QG+ N+ + KED+ L + + V +++T
Sbjct: 313 -----QGRENVKKLLKEDETLRNTIRQQVRDMLT 341
Query= sid | 114832 | 1an | dp1ORF011 Phage dp1 ORF | 28017-29096 | 3
         (359 letters)
>gi|2444110 (U88974) ORF31 [Streptococcus thermophilus temperate bacteriophage
           Length = 348
 Score = 187 bits (469), Expect = 1e-46
Identities = 118/358 (32%), Positives = 187/358 (51%), Gaps = 21/358 (5%)
           {\tt IYDYINAGEIASYIQALPSNALQYLGPTLFPNAQQTGTDISWLKGANNLPVTIQPSNYDA~62}
IYD + A IA Y AL N LG ++FP +Q GT +S++KGA+ V ++ + +D
Sbjct: 4 IYDKVTASNIAGYFNALQENVSSTLGESIFPARKQLGTKLSYIKGASGQSVALKAAAFDT 63
Query: 63 KASLRERAGFSKQATEMAFFRESMRLGEKDRQNLQMLLNQSSA-LAQPLITQLYNDTKNL 121
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+M FF+E+M + E DRQ L ++ + +A L ++ ++ND
 Sbjct: 64 NVTIRDRVSAEMHDEQMPFFKEAMLVKENDRQQLNLVKDSGNAVLVNTIVAGIFNDNLTL 123
 Query: 122 VDGVEAQAEYMRMQLLQYGKFTVKSTNSEAQYTYDYNMDAKQQYAVTKKWTNPAESDPIA 181
V+G A+ E MRMQ+L GK S Y D K+Q V+K W P + P+A Sbjct: 124 VNGARARLEAMRMQVLATGKIAFTSDGVNKDIDYGVKPDHKKQ--VSKSWAEPG-ATPLA 180
 Query: 182 DILAAMDDIENRTGVRPTRMVLNRNTYNQMTKSDSIKKAL-AIGVQGSWENFLLLASDAE 240
           D+ A+ + G+ P R V+N T+ + K+ S K + + GS + ++ E
 Sbjct: 181 DLEDAI-ETARELGLNPERAVMNAKTFGLIRKAASTVKVIKPLAGDGS----AVTKAELE 235
 Query: 241 KFIAEKTGLQIAVYSKKIAQFADADKLPDVGNIRQFNLIDDGKVVLLPPDAVGHTWYGTT 300
                                 D G + +F DG + L+P +G+T +GTT
 Sbjct: 236 NYIADNFGVSIVLENGTYRN------DKGEVSKF--YPDGHLTLIPNGPLGNTVFGTT 285
 Query: 301 PEAFDLASGGT-DAQVQVLSGGPTVTTYLEKHPVNIATVVSAVMIPSFEGIDYVGVLT 357
PE DL + T +A+V+++ G VTT PVN+ T VS V +PSFE +D V +LT Sbjct: 286 PEESDLFADNTVNAEVEIVDNGIAVTTTKTTDPVNVQTKVSMVALPSFERLDDVYMLT 343
Query= sid|114834|lan|dp10RF013 Phage dp1 ORF|10215-11240|3
          (341 letters)
 >sp|P09122|DP3X BACSU DNA POLYMERASE III SUBUNITS GAMMA AND TAU
            Length = 563
 Score = 182 bits (458), Expect = 2e-45
 Identities = 118/353 (33%), Positives = 176/353 (49%), Gaps = 31/353 (8%)
Query: 7 YRPQTFEEVVAQEYVKEILLNQLQNGAIKHGYLFCXXXXXXXXXXXXXXXXXXIFAKDVN----- 60
+RPQ FE+VV QE++ + L N L H YLF +IFAK VN
Sbjct: 10 FRPQRFEDVVGQEHITKTLQNALLQKKFSHAYLFSGPRGTGKTSAAKIFAKAVNCEHAPV 69
Query: 61 ------KGL-----GSPIEIDAASNNGVENVRNIIEDSRYKSMDSEFKVYIIDEVH 105
                     KG+ IEIDAASNNGV+ +R+I + ++ +KVYIIDEVH
Sbjct: 70 DEPCNECAACKGITNGSISDVIEIDAASNNGVDEIRDIRDKVKFAPSAVTYKVYIIDEVH 129
Query: 106 MLSTGAFNALLKTLEEPSSGTVFILCTTDPQKIPDTILSRVQRFDFTRIDNDDIVNQLQF 165
           MLS GAFNALLKTLEEP +FIL TT+P KIP TI+SR QRFDF RI + IV ++
Sbjct: 130 MLSIGAFNALLKTLEEPPEHCIFILATTEPHKIPLTIISRCORFDFKRITSOAIVGRMNK 189
Query: 166 IIESENEEGAGYSYERDALSFIGKLANGGMRDSITRLEKVLDYSHHVDMEAVSNAL---G 222
                        E +L I A+GGMRD+++ L++ + +S D+ V +AL G
Sbjct: 190 IVDAEQ-----LQVEEGSLEIIASAAHGGMRDALSLLDQAISFSG--DILKVEDALLITG 242
Query: 223 VPDYETFASLVEAIANYDGSKCLEIVNDFHYSGKDLKLVTRNFTDFLLEVCKYWLVRDIS 282
                    L +++ + + S LE +N+ GKD + + + ++ Y
Sbjct: 243 AVSQLYIGKLAKSLHDKNVSDALETINELLQQGKDPAKLIEDMIFYFRDMLLYKTAPGLE 302
Query: 283 ITQLPAHFESKLEQFCEAFQYPTLLWMLEEMNELAGVVKWEPNAKPIIETKLL 335
                       + E
                                 L M++ +N+ +KW + + E +-
Sbjct: 303 GVLEKVKVDETFRELSEQIPAQALYEMIDILNKSHQEMKWTNHPRIFFEVAVV 355
Query= sid | 114835 | lan | dp1ORF014 Phage dp1 ORF | 50961-51974 | 3
         (337 letters)
>sp|P47492|PRIM_MYCGE DNA PRIMASE >gi|1361496|pir||F64227 DNA primase (dnaE) homolog
           MG250 - Mycoplasma genitalium (SGC3) >gi|3844848
           (U39704) DNA primase (dnaE) [Mycoplasma genitalium]
           Length = 607
 Score = 57.0 bits (135), Expect = 2e-07
 Identities = 53/190 (27%), Positives = 89/190 (45%), Gaps = 17/190 (8%)
Query: 146 EELDKYRFIHP-----YMYERKLTDELIEMFDVGYDK--LHDCITFPVRNLKGETVFF 196
E +++Y FI+P Y++ K + + FD K + I P+ + G V F
Sbjct: 170 ESMERYPFINPKIKPSELYLFS-KTNQQGLGFFDFNTKKATFQNQIMIPHDFNGNPVGF 228
Query: 197 NRRSVRSKFHQYGEDDPKTEFLYGQYELVAFRDYFEKPISQVFVTESVINCLTLWSMKIP 256 -
+ RSV + ++ EF + + EL+ K ++Q+F+ E + TL + K
Sbjct: 229 SARSVDNINKLKYKNSADHEF-FKKGELLFNFHRLNKNLNQLFIVEGYFDVFTLTNSKFE 287
Query: 257 AVALMGVGGGN-QINLLKR--LPYRNIVLALDPDNAGQTAQEKLYRQLKRSK-VVRFLNY 312
           AVALMG+ + QI +K
                                  + +VLALD D +GQ A L +L + +V + +
Sbjct: 288 AVALMGLALNDVQIKAIKAHFKELQTLVLALDNDASGQNAVFSLIEKLNNNNFIVEIVQW 347
```

Query: 313 PKEFYDNKWD 322

Sbjct: 348 EHNYKD--WD 355

+ D WD

- \_\_\_\_\_

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Query= sid|114837|lan|dp1ORF016 Phage dp1 ORF|43413-44303|3
         (296 letters)
>emb|CAB07986| (Z93946) N-acetylmuramoyl-L-alanine amidase [bacteriophage Dp-1]
           Length = 296
 Score = 661 bits (1686), Expect = 0.0
 Identities = 296/296 (100%), Positives = 296/296 (100%)
          MGVDIEKGVAWMQARKGRVSYSMDFRDGPDSYDCSSSMYYALRSAGASSAGWAVNTEYMH 60
           MGVDIEKGVAWMQARKGRVSYSMDFRDGPDSYDCSSSMYYALRSAGASSAGWAVNTEYMH
Sbjct: 1 MGVDIEKGVAWMQARKGRVSYSMDFRDGPDSYDCSSSMYYALRSAGASSAGWAVNTEYMH 60
Query: 61 AWLIENGYELISENAPWDAKRGDIFIWGRKGASAGAGGHTGMFIDSDNIIHCNYAYDGIS 120
           AWLIENGYELISENAPWDAKRGDIFIWGRKGASAGAGGHTGMFIDSDNIIHCNYAYDGIS
Sbjct: 61 AWLIENGYELISENAPWDAKRGDIFIWGRKGASAGAGGHTGMFIDSDNIIHCNYAYDGIS 120
Query: 121 VNDHDERWYYAGQPYYYVYRLTNANAQPAEKKLGWQKDATGFWYARANGTYPKDEFEYIE 180
           VNDHDERWYYAGQPYYYVYRLTNANAQPAEKKLGWQKDATGFWYARANGTYPKDEFEYIE
Sbjct: 121 VNDHDERWYYAGQPYYYVYRLTNANAQPAEKKLGWQKDATGFWYARANGTYPKDEFEYIE 180
Query: 181 ENKSWFYFDDQGYMLAEKWLKHTDGNWYWFDRDGYMATSWKRIGESWYYFNRDGSMVTGW 240
           ENKSWFYFDDOGYMLAEKWLKHTDGNWYWFDRDGYMATSWKRIGESWYYFNRDGSMVTGW
Sbjct: 181 ENKSWFYFDDQGYMLAEKWLKHTDGNWYWFDRDGYMATSWKRIGESWYYFNRDGSMVTGW 240
Query: 241 IKYYDNWYYCDATNGDMKSNAFIRYNDGWYLLLPDGRLADKPQPTVEPDGLITAKV 296
           IKYYDNWYYCDATNGDMKSNAFIRYNDGWYLLLPDGRLADKPQFTVEPDGLITAKV
Sbjct: 241 IKYYDNWYYCDATNGDMKSNAFIRYNDGWYLLLPDGRLADKPQFTVEPDGLITAKV 296
Query= sid|114841|lan|dp1ORF020 Phage dp1 ORF|1864-2658|1
         (264 letters)
>emb|CAB13247| (299111) similar to coenzyme PQQ synthesis [Bacillus subtilis]
          Length = 243
 Score = 217 bits (548), Expect = 5e-56
 Identities = 117/248 (47%), Positives = 163/248 (65%), Gaps = 15/248 (6%)
Ouery: 23 MPIMEIFGPTIOGEGMVIGOKTIFIRTGGCDYHCNWCDSAFTWNGTTEPE--YITGKEAA 80
           +P++EIFGPTIQGEGMVIGQKT+F+RT GCDY C+WCDSAFTW+G+ + + ++T +E
          IPVLEIFGPTIOGEGMVIGOKTMFVRTAGCDYSCSWCDSAFTWDGSAKKDIRWMTAEEIF 64
Query: 81 SRILKLAFNDKGEQICNHVTLTGGNPALINEPMAKMISILKEHGFKFGLETQGTRFQEWF 140
                         +HVT++GGNPAL+ + + I +LKE+ + LETQGT +Q+WF
                 DG
Sbjct: 65 AEL----KDIGGDAFSHVTISGGNPALLKQ-LDAFIELLKENNIRAALETQGTVYQDWF 118
Query: 141 KEVSDITISPKPPSSGMRTNMKILEAIVDRM--NDENLDWSFKIVIFDENDLAYARDMFK 198
             + D+TISPKPPSS M TN + L+ I+ + ND
                                                 S K+VIF++ DL +A+ + K
Sbjct: 119 TLIDDLTISPKPPSSKMVTNFQKLDHILTSLQENDRQHAVSLKVVIFNDEDLEFAKTVHK 178
Query: 199 TFEGKLRPVNYLSVGNANAY--EEGKISDRLLEKLGWLWDKVYEDPAFNNVRPLPQLHTL 256
                   YL VGN + + ++ + LL K L DKV D N VR LPQLHTL
Sbjct: 179 RYPG---IPFYLQVGNDDVHTTDDQSLIAHLLGKYEALVDKVAVDAELNLVRVLPQLHTL 235
Query: 257 VYDNKRGV 264
           ++ NKRGV
Sbjct: 236 LWGNKRGV 243
```

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Query= sid|114842|lan|dp1ORF021 Phage dp1 ORF|2504-3295|2
         (263 letters)
>sp|P19465|GCH1 BACSU GTP CYCLOHYDROLASE I (GTP-CH-I) >gi|98411|pir||A38256 GTP
           cyclohydrolase I (EC 3.5.4.16) - Bacillus subtilis
           >gi|143231 (M37320) regulatory protein (Bacillus
           subtilis) >gi|143799 (M80245) MtrA (Bacillus subtilis)
           >gi|2634696|emb|CAB14194| (Z99115) GTP cyclohydrolase I
            [Bacillus subtilis]
           Length = 190
 Score = 208 bits (523), Expect = 4e-53
 Identities = 103/185 (55%), Positives = 133/185 (71%), Gaps = 1/185 (0%)
Query: 80 VTLDNTEAAVQRLFGLLGEDAERDGLQDTPFRFVKALAEHTVGYREDPKLHLEKTFDVDH 139
           V + E AV+++ +GED R+GL DTP R K AE G EDPK H + F
           VNKEQIEQAVRQILEAIGEDPNREGLLDTPKRVAKMYAEVFSGLNEDPKEHFQTIFGENH 63
Query: 140 EDLVLVKDIPFNSLCEHHLAPFVGKVHIAYIPKD-KITGLSKFGRVVEGYAKRLQVQERL 198
           E+LVLVKDI F+S+CEHHL PF GK H+AYIP+ K+TGLSK R VE AKR Q+QER+
Sbjct: 64 EELVLVKDIAFHSMCEHHLVPFYGKAHVAYIPRGGKVTGLSKLARAVEAVAKRPQLQERI 123
Query: 199 TQQIADAIQEVLNPQAVAVIVEAEHTCMSGRGIKKHGATTVTSTMRGLFQDDASARAELL 258
            T IA++I E L+P V V+VEAEH CM+ RG++K GA TVTS +RG+F+DDA+ARAE+L
Sbjct: 124 TSTIAESIVETLDPHGVMVVVEAEHMCMTMRGVRKPGAKTVTSAVRGVFKDDAAARAEVL 183
Query: 259 QLIKK 263
Sbjct: 184 EHIKR 188
Query= sid|114843|lan|dplORF022 Phage dp1 ORF|30896-31675|2
         (259 letters)
>gi|2347102 (U77367) internalin [Listeria monocytogenes]
           Length = 821
 Score = 55.0 bits (130). Expect = 5e-07
 Identities = 44/149 (29%), Positives = 63/149 (41%), Gaps = 13/149 (8%)
Query: 119 FRMNIYVPNYVG--DSIVNYVKITLNNCTGKAPGLSIGKEFYAPEFNIKAREATKAGLPV 176
             + VPN + D + + NN T AP L
                                                     Y PE +K + K
Sbjct: 383 FSKTLSVPNNITSIDGTLIAPETISNNGTYDAPNLKWSLPNYLPE--VKYTFSQKIPIGT 440
Query: 177 KSMDYVAQLPAVLR-----RVTFDLNGGTGTADAVRVEAGKKISPKPVDPTLTGKAFKGW 231
+ +Y + L+ +VTF++ G T + V E + P+P PT G F GW
Sbjct: 441 GTSNYSGFITQPLKELLDYKVTFNVEGNTSEVETVTEE---NLIPEPTSPTKQGYTFDGW 497
Query: 232 -KVEGESTIWDFDNHMMPDRDVKLVAQFA 259
              E T WDF MP D+ L A F+
Sbjct: 498 YDAETGGTKWDFTTGQMPANDLTLYAHFS 526
 Score = 43.4 bits (100), Expect = 0.002
 Identities = 47/195 (24%), Positives = 73/195 (37%), Gaps = 12/195 (6%)
Query: 72 YDLTFKDNTFDPEIMALIEGGTVRQQGGTIAGYDT-PMLAQGASNMKPFRMNIYVPNY-- 128
YD + T + +G + GG + T M A + F +N Y N+
Sbjct: 547 YDALLNEPTTPTKQGYTFDGWYDAETGGNKWDFKTMKMPANDVAFYAHFTINNYQANFDI 606
Query: 129 ---VGDSIVNYVKITLNNCTGKAPGLSIGKEFYAPEFNIKAREATKAGLPVKSMDYVAQL 185
              V + + Y + T G + + A K TK +P +
Sbjct: 607 DGEVKNETIAYDTLLNEPTTPTKQGYTFDGWYDAETGGTKWDFKTKE-MPANDVTLYAHF 665
Query: 186 PAVLRRVTFDLNGGTGTADAVRVEAGKKISPKPVDPTLTGKAFKGW-KVEGESTIWDFDN 244
+ FD++G T + V +A + P+P P+ TG +GW E T WDF
Sbjct: 666 TINNYQANFDIDGAV-TEEVVNYDA---LIPEPTSPSKTGFTLEGWYDAEVGGTKWDFKT 721
Query: 245 HMMPDRDVKLVAQFA 259
                                                                                 - _____
             MP D+ L A F+
Sbjct: 722 MKMPANDITLYAHFS 736
 Score = 38.3 bits (87), Expect = 0.057
 Identities = 42/169 (24%), Positives = 59/169 (34%), Gaps = 10/169 (5%)
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Query: 96 QQGGTIAGYDT-PMLAQGASNMKPPRMNIYVPNYVGDSIVNYVKIT----LNNCTGKAPG 150
   + GGT + T M A + F +N Y N+ D +V + LN T
Sbjct: 501 ETGGTKWDFTTGQMPANDLTLYAHFSVNSYQANFDIDGVVTNEAVVYDALLNEPTTPTKQ 560
   Query: 151 LSIGKEFYAPEFNIKAREATKAGLPVKSMDYVAQLPAVLRRVTFDLNGGTGTADAVRVEA 210
                      +Y E
                                 + +P ++ A
                                                         + FD++G
   Sbjct: 561 GYTFDGWYDAETGGNKWDFKTMKMPANDVAFYAHFTINNYQANFDIDGEVKNETI----A 616
   Query: 211 GKKISPKPVDPTLTGKAFKGW-KVEGESTIWDFDNHMMPDRDVKLVAQF 258
                   + +P PT G F GW E T WDF MP DV LAP
   Sbjct: 617 YDTLLNEPTTPTKQGYTFDGWYDAETGGTKWDFKTKEMPANDVTLYAHF 665
   Query= sid|114850|lan|dp10RF029 Phage dp1 ORF|662-1348|2
             (228 letters)
   >gi|2650185 (AE001074) succinoglycan biosynthesis regulator (exsB)
               [Archaeoglobus fulgidus]
               Length = 239
    Score = 119 bits (295), Expect = 2e-26 Identities = 79/224 (35%), Positives = 113/224 (50%), Gaps = 11/224 (4%)
   Query: 1 MKSVVLLSGGVDSATCLAIEVDKWGSKNVHAIAFNYGQKHEAELENAANVAMFYGVKFTI 60
   MK-V+LLSGG+DS+T L +D G VHA+ F YGQKH E+E+A VA V+
Sbjct: 1 MKAVMLLSGGIDSSTLLYYLLD--GGYEVHALTFFYGQKHSKEIESAEKVAKAAKVRHLK 58
   Query: 61 LEIDSKIYXXXXXXLLQGKGEISHGKSYAEILAEKEVVDTYVPFRNGLMLSQXXXXXXXX 120
   ++I S I+ L G+ E+ Y+E + + T VP RN ++LS
Sbjct: 59 VDI-STIHDLISYGALTGEBEVPKA-FYSEEVQRR----TIVPNRNMILLS--IAAGYAV 110
   Query: 121 XXXXXXXXXXXXXXXXXXXXXPDCTPEFYNSMSNAMEYGT-GGKVTLVAPLLTLTKAQVVKW 179
PDC EF ++ A+ V + AP + +TKA +V+
Sbjct: 111 KIGAKEVHYAAHLSDYSIYPDCRKEFVKALDTAVYLANIWTPVEVRAPFVDMTKADIVRL 170
   Query: 180 GIDLDVPYFLTRSCYESDAESCGTCATCIDRKKAFEENGMTDPI 223
   G+ L VPY LT SCYE C +C TC++R +AF NG+ DP+
Sbjct: 171 GLKLGVPYELTWSCYEGGDRPCLSCGTCLERTEAFLANGVKDPL 214
   Query= sid|114855|lan|dp1ORF034 Phage dp1 ORF|131-652|2
             (173 letters)
   >emb|CAB13248| (Z99111) similar to hypothetical proteins [Bacillus subtilis]
               Length = 165
    Score = 220 bits (556), Expect = 4e-57
    Identities = 103/139 (74%), Positives = 117/139 (84%)
   Query: 5 TTRTDAELTGVTLLGNQDTKYDYDYNPDVLETFPNKHPENNYLVTFDGYEFTSLCPKTGQ 64
  TTR ++EL GVTLLGNQ T Y ++Y PDVLE+FPNKH +Y V F+ EFTSLCPKTGQ
Sbjct: 2 TTRKESELEGVTLLGNQGTNYLFEYAPDVLESFPNKHVNRDYFVKFNCPEFTSLCPKTGQ 61
   Query: 65 PDFANVPISYIPNEKMVESKSLKLYLFSFRNHGDFHEDCMNIILNDLYELMEPKYIEVMG 124
               PDFA ++ISYIP+EKMVESKSLKLYLFSFRNHGDFHEDCMNII+NDL ELM+P+YIEV G
   Sbjct: 62 PDFATIYISYIPDEKMVESKSLKLYLFSFRNHGDFHEDCMNIIMNDLIELMDPRYIEVWG 121
   Query: 125 LFTPRGGISIYPFVNKVNP 143
                PTPRGGISI P+ N
   Sbjct: 122 KFTPRGGISIDPYTNYGKP 140
   Query= sid|114857|lan|dp10RF036 Phage dp1 ORF|48808-49362|1
             (184 letters)
   >gi|1353529 (U38906) ORF12 [Bacteriophage r1t]
               Length = 296
    Score = 53.5 bits (126), Expect = 1e-06
    Identities = 42/149 (28%), Positives = 70/149 (46%), Gaps = 9/149 (6%)
   Query: 34 IASNTVGNGKTSWAVRLLQRYLAETALDGRIVEKGMFVVSAQLLTEFGDYNYFQTMQEFL 93
   + S G GK+ A+ +L+ L T L ++ V + F + + F + + F +
Sbjct: 155 VVSGPAGTGKSHLAMSILKDCLQHTDLT--VIFASWSEVLHLIKDSFDNKDSFYSTEYFM 212
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Query: 94 ERFERLKTCELLVIDEIGGGSLTKASYPYLYDLVNYRVDNNLSTIYTTNYTDDEIIDLLG 153
           E F + +LLVID+IG +T+ S L ++++ R
                                                     TI TTN DEI
Sbjct: 213 EVF---RNTDLLVIDDIGSEKITEWSMSLLTEVLDART----KTIITTNLKSDEIRKKYH 265
Query: 154 QRLYSRIYDTSVVLDFQASNVRGLEVSEI 182
            R YSR++
                         F N++
Sbjct: 266 NRTYSRLFRGIGKKAFNFENIKDKRVSQL 294
Query= sid|114859|lan|dp10RF038 Phage dp1 ORF|1350-1871|3
         (173 letters)
>8p|P44123|YB90_HAEIN HYPOTHETICAL PROTEIN HI1190 >gi|1074675|pir||F64021 hypothetical
           protein HI1190 - Haemophilus influenzae (strain Rd KW20)
           >gi|1574117 (U32798) 6-pyruvoyl tetrahydrobiopterin
           synthase, putative [Haemophilus influenzae Rd]
           Length = 141
 Score = 100 bits (247), Expect = 6e-21
 Identities = 59/143 (41%), Positives = 83/143 (57%), Gaps = 10/143 (6%)
Query: 2 RVSKTLTFDAAHQLVGHFGKCANLHGHTYKVEISLAGGTYDHGSSQGMVVDFYHVKKIA- 60
++SK +FD AH L GH GKC NLHGHTYK+++ ++G Y G+ + MV+DF +K I
Sbjct: 3 KISKEFSFDMAHLLDGHDGKCQNLHGHTYKLQVEISGDLYKSGAKKAMVIDFSDLKSIVK 62
Query: 61 GTFIDRLDHAVLL-QGNEP----IALANAVDTKRVLFGFRTTAENMSRFLTWTLTELMWK 115
+D +DHA + Q NE L +++K FRTTAE ++RF+ L +
Sbjct: 63 KVILDPMDHAFIYDQTNERESQIATLLQKLNSKTFGVPFRTTAEBIARFIFNRLKH--DB 120
Query: 116 HARIDSIKLWETPTGCAECTYYE 138
I SI+LWETPT + C Y E
Sbjct: 121 QLSISSIRLWETPT--SFCEYQE 141
Query= sid|114860|lan|dp1ORF039 Phage dp1 ORF|3306-3803|3
         (165 letters)
>emb|CAA68244| (X99978) ORF7; hydophobic protein [Lactobacillus plantarum]
           Length = 168
 Score = 64.4 bits (154), Expect = 5e-10
 Identities = 49/156 (31%), Positives = 84/156 (53%), Gaps = 9/156 (5%)
Query: 8 WLVRTALIAALYVTLTVAFSAISY--GPIQFRVSEALILLPLWNHRWTPGIVLGTIIANF 65
           W++ AL+AA+YV L + +A S G IQFRVSE L L ++N ++ GIV G I+ +
Sbjct: 9
           WIIN-ALVAAMYVVLCLGPAAFSLASGAIQFRVSEGLNHLAVFNRKYIWGIVAGVILFDA 67
Query: 66 FSP-LGLIDVLFGSLATFLGXXXXXXXXXXXXSPLYSLICPVLA----NAYLIALELRIVY 120
           F P L++VLFG + L
                                             ++ + +A + ++IAL + ++
Sbjct: 68 FGPGASLLNVLFGGGQSLLALLVLTWLAPKLKTVWQRMLLNIALFTVSMFMIALMITMMS 127
Query: 121 S-LPFWESVIYVGISEAIIVLISYFLISTLAKNNHF 155
S + FW + + +SE II+ I+ ++ +L + HF
Sbjct: 128 SGVAFWPTYLTTALSELIIMSITAPIMYSLDRVLHF 163
Query= sid|114862|lan|dp1ORF041 Phage dp1 ORF|8208-8699|3
         (163 letters)
>gi|2522313 (AF012906) dUTPase homolog [Bacillus subtilis]
           >gi|2634394|emb|CAB13893| (Z99114) similar to
           deoxyuridine 5'-triphosphate nucleotidohydrolase
           [Bacillus subtilis] >gi|3025643 (AF020713) putative
           dUTPase (Bacteriophage SPBc2)
           Length = 142
 Score = 108 bits (267), Expect = 2e-23
 Identities = 65/160 (40%), Positives = 83/160 (51%), Gaps = 25/160 (15%)
           VDVKMIDPKLDRLKYT--GDWVDVRISSITKIDADSADVSRCRKVLQKAQVYSVAAGECI 62
Query: 5
           + +K +D R+ GDW+D+R + I D +
           IKIKYLDETQTRINKMEQGDWIDLRAAEDVAIKKDEFKL------41
Sbjct: 3
Query: 63 KIAHGFALELPKGYEAILHPRSSLFKKTGLIFVSS-GVIDEGYKGDTDEWFSVWYATRDA 121
              G A+ELP+GYEA + PRSS +K G+I +S GVIDE YKGD D WF YA RD
Sbjct: 42 -VPLGVAMELPEGYEAHVVPRSSTYKNFGVIQTNSMGVIDESYKGDNDFWFFPAYALRDT 100
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WO 00/32825 PCT/IB99/02040

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Query: 122 DIFYDQRIAQFRIQEKQPAIKFNFVESLGNAARGGHGSTG 161
I RI QFRI +K PA+ V+ LGN RGGHGSTG
Sbjct: 101 KIKKGDRICQFRIMKKMPAVDLIEVDRLGNGDRGGHGSTG 140
Query= sid|114867|lan|dplORF046 Phage dpl ORF|42774-43202|3
         (142 letters)
>emb|CAB07984| (293946) hypothetical protein [bacteriophage Dp-1]
 Score = 287 bits (728), Expect = 2e-77
 Identities = 142/142 (100%), Positives = 142/142 (100%)
           MPMWLNDTAVLTTIITACSGVLTVLLNKLFEWKSNKAKSVLEDISTTLSTLKQQVDGIDQ 60
            MPMWLNDTAVLTTI ITACSGVLTVLLNKLFEWKSNKAKSVLEDISTTLSTLKQQVDGIDQ
           MPMWLNDTAVLTTIITACSGVLTVLLNKLFEWKSNKAKSVLEDISTTLSTLKQQVDGIDQ 60
 Query: 61 TTVAINHQNDVIQDGTRKIQRYRLYHDLKREVITGYTTLDHFRELSILFESYKNLGGNGE 120
            TTVAINHQNDVIQDGTRKIQRYRLYHDLKREVITGYTTLDHFRELSILFESYKNLGGNGE
 Sbjct: 61 TTVAINHQNDVIQDGTRKIQRYRLYHDLKREVITGYTTLDHFRELSILFESYKNLGGNGE 120
 Query: 121 VEALYEKYKKLPIREEDLDETI 142
            VEALYEKYKKLPIREEDLDETI
 Sbjct: 121 VEALYEKYKKLPIREEDLDETI 142
 Query= sid|114901|lan|dp10RF080 Phage dp1 ORF|42490-42759|1
          (89 letters)
 >emb|CAB07983| (Z93946) hypothetical protein [bacteriophage Dp-1]
           Length = 124
  Score = 147 bits (367), Expect = 1e-35
  Identities = 75/75 (100%), Positives = 75/75 (100%)
 Query: 1 MLNLTKSRQIVAEFTIGQGAEKKLVKTTIVNIDANAVSTVSETLHDPDLYAANRRELRAD 60
           MLNLTKSRQIVAEFTIGQGAEKKLVKTTIVNIDANAVSTVSETLHDPDLYAANRRELRAD
 Sbjct: 1 MLNLTKSRQIVAEFTIGQGAEKKLVKTTIVNIDANAVSTVSETLHDPDLYAANRRELRAD 60
 Query: 61 EQKLRETRYAIEDEI 75
           EQKLRETRYALEDEI
  Sbjct: 61 EQKLRETRYAIEDEI 75
  Query= sid|114912|lan|dp1ORF091 Phage dp1 ORF|43189-43413|1
           (74 letters)
  >emb|CAB07985| (Z93946) holin [bacteriophage Dp-1]
            Length = 74
   Score = 63.2 bits (151), Expect = 2e-10
   Identities = 34/74 (45%), Positives = 34/74 (45%)
  VLGVSSR
                                            YQFD
  Sbjct: 1 MKLSNEQYDVAKNVVTVVVPAAIALITGLGALYQFDTTAITGTIALLATFAGTVLGVSSR 60
            MKLSNEQYD
  Query: 61 NYQKEQEAQNNEVE 74
            NYOKEQEAQNNEVE
  Sbjct: 61 NYQKEQEAQNNEVE 74
```

## Condensed listing of homology information from above

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Phage: dpl
Database: nr
Program: Blastp
Query= sid|114822|lan|dp10RF001 Phage dp1 ORF|36698-40390|2
          (1230 letters)
gi|2444124 (U88974) ORF45 (Streptococcus thermophilus temperate ...
gi|928828 (L44593) ORF1904; putative [Lactococcus lactis phage B...
                                                                                  e-118
                                                                             427
gi|2935676 (AF032121) unknown [Streptococcus thermophilus bacter...
                                                                                  1e-82
                                                                             309
gi|2935691 (AF032122) unknown (Streptococcus thermophilus bacter...
                                                                             306
                                                                                  7e-82
gi|3540289 (AF057033) putative anti-receptor [Streptococcus ther...
                                                                             279
                                                                                   6e-74
gi|4530154|gb|AAD21894.1| (AF085222) putative tail-host specific...
                                                                                  3e-56
                                                                             220
gi|930045|emb|CAA33387| (X15332) alpha-1 (III) collagen [Homo sa...
                                                                                   4e-07
gi|1070603|pir||CGHU7L collagen alpha 1(III) chain precursor - h...
                                                                              58
                                                                                   4e-07
gi|4502951|ref|NP_000081.1|PCOL3A1| collagen, type III, alpha 1 ...
                                                                                   4e-07
gi|115290|sp|P04258|CA13_BOVIN COLLAGEN ALPHA 1(III) CHAIN >gi|7...
                                                                              58
                                                                                   4e-07
gi|575322|emb|CAA36279| (X52046) type III collagen [Mus musculus]
                                                                              57
                                                                                   8e-07
gi|2119163|pir||S59856 collagen alpha 1(III) chain precursor - m...
                                                                              57
                                                                                   8e-07
gi|543912|sp|P13941|CA13_RAT COLLAGEN ALPHA (IIII) CHAIN >gi|543...
gi|3171998|emb|CAA06510| (AJ005395) collagen alpha 1 (III) [Ratt...
gi|3947565|emb|CAA90250| (Z49967) similar to collagen; cDNA EST ...
                                                                              57
                                                                                   1e-06
                                                                              57
                                                                                   le-06
                                                                              54
                                                                                   7e-06
gi|423403|pir||A46053 bullous pemphigoid antigen, BPAG2, type XV...
                                                                               53
                                                                                   9e-06
gi|115410|sp|P12114|CCS1_CAEEL_CUTICLE_COLLAGEN_SQT-1 >gi|84437|...
gi|3873801|emb|CAA90084| (Z49907) cuticle_collagen_SQT-1; cDNA_E...
                                                                               53
                                                                                   9e-06
                                                                                   9e-06
Query= sid|114823|lan|dp10RF002 Phage dp1 ORF|32386-35835|1
          (1149 letters)
gi|3341922|dbj|BAA31888| (AB009866) orf 15 [bacteriophage phi PVL]
                                                                             280 3e-74
gi|4126622|dbj|BAA36642.1| (AB016282) ORF36 [bacteriophage phi-105]
                                                                                   1e-59
                                                                              232
gi|1369948|emb|CAA59194| (X84706) host interacting protein [Bact...
                                                                                   3e-50
                                                                              201
                                                                                   2e-46
gi|3139112 (AF063097) gpT [Bacteriophage P2]
                                                                              188
gi 3337272 (U32222) G protein [Bacteriophage 186]
                                                                             161
gi|4063799|dbj|BAA36253| (AB008550) orf25; similar to T gene of ...
                                                                              159
                                                                                   8e-38
gi|3172274 (AF022214) minor tail subunit; putative tape-measure ...
                                                                              123
                                                                                   6e-27
gi|465127|sp|Q05233|VG26_BPML5 MINOR TAIL PROTEIN GP26 >gi|41904...
                                                                              108
                                                                                   2e-22
gi 3540284 (AF057033) putative minor tail protein [Streptococcus...
gi|2444119 (U88974) ORF40 (Streptococcus thermophilus temperate ...
                                                                                   6e-17
gi|2634555|emb|CAB14053| (Z99115) yomI [Bacillus subtilis] >gi|3...
                                                                                   1e-09
gi|2392838 (AF011378) unknown [Bacteriophage skl]
                                                                               64
                                                                                   5e-09
                                                                                   3e-08
gi|2764873|emb|CAA66557| (X97918) gene 18.1 [Bacteriophage SPP1]
                                                                                   6e-08
gi|1353559 (U38906) ORF42 [Bacteriophage rlt]
                                                                               61
gi|630841|pir||S39079 puff C-8 protein - fungus gnat (Rhynchosci...
                                                                                   2e-06
gi 1730865 sp P51731 Y027 BPHP1 HYPOTHETICAL 72.8 KD PROTEIN IN ...
                                                                               53
                                                                                   8e-06
gi|224288|prf||1101273J ORF 7 [Bacteriophage HP1]
                                                                                   1e-05
 Query= sid|114824|lan|dp10RF003 Phage dp1 ORF|53538-55877|3
           (779 letters)
 gi|118825|sp|P00582|DP01_ECOLI DNA POLYMERASE I (POL I) >gi|6705...
                                                                              193 3e-48
 gi|2982102|pdb|1KFS|A Chain A, All-Oxygen Dna Complexed To The 3...
                                                                              193
                                                                                   3e-48
                                                                                   3e-48
gi|229889|pdb|1DPI| DNA Polymerase I (Klenow Fragment) (E.C.2....
gi|1169402|sp|P43741|DPOI_HAEIN DNA POLYMERASE I (POL I) >gi|107...
                        DNA Polymerase I (Klenow Fragment) (E.C.2....
                                                                              193
                                                                              191
                                                                                   1e-47
 gi|2688462 (AE001156) DNA polymerase I (polA) [Borrelia burgdorf...
                                                                              190
                                                                                   3e-47
                                                                                   3e-47
 gi|809180|pdb|1KLN|A Escherichia coli
                                                                              190
 gi|1913934|emb|CAA72997| (Y12328) DNA-directed DNA polymerase I ...
                                                                                   8e-47
                                                                              189
 gi|4090935 (AF028719) DNA polymerase type I (Rhodothermus sp. 'I...
                                                                              175
                                                                                   1e-42
 gi|4731571|gb|AAD28505.1|AF121780_1 (AF121780) DNA polymerase I ...
                                                                              174
                                                                                   2e-42
 gi 1633576 (US7757) similar to proofreading 3'-5' exonuclease an...
                                                                              173
                                                                                   4e-42
 gi|3322368 (AE001195) DNA polymerase I (polA) [Treponema pallidum]
                                                                              172
 gi|1006595|dbj|BAA10748| (D64005) DNA polymerase I [Synechocysti...
                                                                              171
 gi|585062|sp|Q07700|DPO1_MYCTU DNA POLYMERASE I (POL I) >gi|4161...
                                                                                   5e-39
 gi|4376908|gb|AAD18751| (AE001645) DNA Polymerase I [Chlamydia p...
                                                                              157
 gi|1169403|sp|P46835|DP01_MYCLE DNA POLYMERASE I (POL I) >gi|107...
 gi 2145839 pir | S72949 DNA polymerase I - Mycobacterium leprae >...
gi 1405438 emb CAA67184 | (X98575) DNA-dependent DNA polymerase [...
                                                                              152
                                                                                    7e-36
                                                                                   9e-36
 gi 2506365 sp P80194 DP01_THECA DNA POLYMERASE I, THERMOSTABLE (...
                                                                                   2e-34
                                                                              147
 gi|3328929 (AE001322) DNA Polymerase I [Chlamydia trachomatis]
                                                                              147 3e-34
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gi|3913510|sp|052225|DP01_THEFI DNA POLYMERASE I, THERMOSTABLE (...
                                                                                     146
                                                                                           7e-34
gi 1205984 (U33536) DNA polymerase I [Bacillus stearothermophilus]
                                                                                           7e-34
gi|118827|sp|P13252|DP01_STRPN DNA POLYMERASE I (POL I) >gi|9802...
                                                                                     145
                                                                                           9e-34
gi|1942202|pdb|1JXE| Stoffel Fragment Of Taq Dna Polymerase I
gi|1943520|pdb|1KTQ| Dna Polymerase
                                                                                           le-33
                                                                                     145
                                                                                           1e-33
gi 1084022 pir | JX0359 DNA-directed DNA polymerase (EC 2.7.7.7) ...
                                                                                           1e-33
                                                                                     145
gi|507891|dbj|BAA06775| (D32013) DNA Polymerase [Thermus aquaticus]
                                                                                     145
                                                                                           le-33
gi 118828 sp P19821 DPO1_THEAQ DNA POLYMERASE I, THERMOSTABLE (T...
                                                                                     145
                                                                                           le-33
gi|1706502|sp|P52028|DP01_THETH DNA POLYMERASE I, THERMOSTABLE (...
                                                                                     144
                                                                                           2e-33
gi|1097211|prf||2113329A DNA polymerase [Thermus aquaticus therm...
                                                                                     144
                                                                                           2e-33
gi 2098289 pdb 1TAU A Chain A, Structure Of Dna Polymerase
                                                                                     143 3e-33
Query= sid | 114825 | lan | dp10RF004 Phage dp1 ORF | 40401-42440 | 3
           (679 letters)
gi|1934761|emb|CAB07981| (Z93946) hypothetical protein (bacterio...
gi|3540290 (AF057033) putative minor structural protein [Strepto...
                                                                                           2e-94
gi|2444125 (U88974) ORF46 (Streptococcus thermophilus temperate ...
                                                                                           3e-92
gi|1934762|emb|CAB07982| (293946) hypothetical protein (bacterio...
                                                                                     300
                                                                                           2e-80
gi|4530155|gb|AAD21895.1| (AF085222) unknown [Streptococcus ther...
                                                                                     276
                                                                                           4e-73
gi 2935677 (AF032121) unknown (Streptococcus thermophilus bacter... gi 2935692 (AF032122) unknown (Streptococcus thermophilus bacter...
                                                                                     250
                                                                                           3e-65
gi 1136289 (U42597) histidine kinase A [Dictyostelium discoideum]
Query= sid|114827|lan|dp10RF006 Phage dp1 ORF|45296-46987|2
           (563 letters)
gi|4377165|gb|AAD18987| (AE001666) SWI/SNF family helicase_2 [Ch...
gi|1769947|emb|CAA67095| (X98455) SNF [Bacillus cereus]
gi|3329163 (AE001341) SWF/SNF family helicase [Chlamydia trachom...
gi|4377149|gb|AAD18973| (AE001664) SWI/SNF family helicase_1 [Ch...
                                                                                     171 le-41
                                                                                     160
                                                                                           3e-38
                                                                                     159
                                                                                           6e-38
                                                                                     157
                                                                                           2e-37
gi 3328995 (AE001326) SWI/SNF family helicase [Chlamydia trachom...
                                                                                     153
                                                                                           2e-36
gi 2493354 sp P75093 Y018 MYCPN HYPOTHETICAL HELICASE MG018/MG01...
                                                                                     146
                                                                                           4e-34
gi | 1653748 | db | | BAA18659 | (D90916) helicase of the snf2/rad54 fam...
gi | 1763712 | emb | CAB05939 | (283337) member of the SNF2 helicase fa...
                                                                                     143
                                                                                           3e-33
                                                                                     143
                                                                                           4e-33
gi|2636153|emb|CAB15645.1| (299122) similar to SNF2 helicase [Ba...
gi|2909552|emb|CAA17284| (AL021924) helZ [Mycobacterium tubercul...
                                                                                     143
                                                                                           4e-33
                                                                                     140
                                                                                           2e-32
gi|3844627 (U39681) ATP-dependent RNA helicase, putative (Mycopl...
gi|3351463|sp|P47264|Y018_MYCGE HYPOTHETICAL HELICASE MG018
                                                                                    136
                                                                                          3e-31
                                                                                     136
                                                                                           46-31
gi 2660669 (AC002342) human Mi-2 autoantigen-like protein [Arabi...
                                                                                     131
                                                                                           2e-29
gi|1361537|pir||164201 helicase (mot1) homolog - Mycoplasma geni...
                                                                                     129
                                                                                           4e-29
gi|3482977|emb|CAA20533.1| (AL031369) putative protein [Arabidop...
gi|3298562 (U91543) zinc-finger helicase [Homo sapiens]
                                                                                     128
                                                                                           9e-29
                                                                                     120
                                                                                           2e-26
gi 3875971 emb CAB02491 (Z80344) similar to helicase; cDNA EST ...
                                                                                     120
                                                                                           2e-26
gi|4557451|ref|NP_001263.1|PCHD3| chromodomain helicase DNA bind...
                                                                                    120 2e-26
gi 2645435 (AF007780) CHD3 (Drosophila melanogaster)
                                                                                     118
                                                                                          le-25
gi|3875165|emb|CAA91798| (Z67881) Similarity to Mouse Chromodoma...
                                                                                          1e-25
Query= sid|114828|lan|dp10RF007 Phage dp1 ORF|22230-23621|3
           (463 letters)
gi|2444105 (U88974) ORF26 (Streptococcus thermophilus temperate ...
                                                                                      89 7e-17
gi|3318666 (U19754) BBA31 homolog [Borrelia burgdorferi]
                                                                                      59
                                                                                          7e-08
gi|2690260 (AE000790) conserved hypothetical protein [Borrelia b...
                                                                                      56
                                                                                          5e-07
Query= sid|114829|lan|dp1ORF008 Phage dp1 ORF|49624-50961|1
           (445 letters)
gi|4406210|gb|AAD19901| (AF100420) DnaB replication fork helicas...
                                                                                      68
                                                                                          2e-10
gi 3121983 sp 025916 DNAB HELPY REPLICATIVE DNA HELICASE >gi 231...
                                                                                      67
                                                                                          2e-10
gi|4416322|gb|AAD20314| (AF106032) replicative helicase; DnaB [B...
                                                                                      65
                                                                                          9e-10
gi 4155895 (AE001551) REPLICATIVE DNA HELICASE (Helicobacter pyl...
                                                                                      60
                                                                                          4e-08
gi|3322317 (AE001191) replicative DNA helicase (dnaB) [Treponema...
gi|138031|sp|P04530|VG41_BPT4 PRIMASE-HELICASE (PROTEIN GP41) >g...
                                                                                      58
                                                                                          le-07
                                                                                      53
                                                                                          3e-06
gi 2983861 (AE000742) replicative DNA helicase [Aquifex aeolicus]
                                                                                      51 1e-05
Query= sid|114831|lan|dp10RF010 Phage dp1 ORF|8699-9859|2
           (386 letters)
                                                                                                     _____
gi|2760912 (AF037258) RecA protein [Chlorobium tepidum]
                                                                                          2e-30 _
                                                                                    133
gi|3219851|sp|P94666|RECA_CLOPE RECA PROTEIN >gi|1698591 (U61497...
                                                                                    129
                                                                                          3e-29
gi|1350566|sp|P48295|RECA_STRVL RECA PROTEIN >gi|508860 (U04837)...
                                                                                    128
                                                                                          7e-29
gi|744163|prf||2014250A recA-like protein (Streptomyces violaceus)
                                                                                    126
                                                                                          3e-28
gi|730487|sp|P41054|RECA_STRAM_RECA_PROTEIN >gi|511133|emb|CAAA2...
gi|2687334|emb|CAA15875| (AL020958) RecA_protein [Streptomyces c...
gi|1350565|sp|P48294|RECA_STRLI_RECA_PROTEIN >gi|481482|pir||S38...
                                                                                    125
                                                                                          4e-28
                                                                                    125
                                                                                          6e-28
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gi|464599|sp|P33542|RECA_AQUPY RECA PROTEIN >gi|1086167|pir||A55...
                                                                                         123 2e-27
gi|417636|sp|P32725|RECA_RHOSH RECA PROTEIN >gi|541307|pir||S415...
                                                                                         123 2e-27

      gi|2984348 (AE000775)
      recombination protein RecA [Aquifex aeolicus]

      gi|3219854
      lsp|P95846 [RECA STRRM RECA PROTEIN >gi|1729800 | emb|CAA...

      gi|2500086
      sp|Q59560 | RECA MYCSM RECA PROTEIN >gi|1430892 | emb|CAA...

      gi|1350567
      sp|P48296 | RECA THEAQ RECA PROTEIN >gi|1072963 | pir | A5...

                                                                                                2e-27
                                                                                         123
                                                                                                4e-27
                                                                                         122
                                                                                                4e-27
                                                                                         122
                                                                                         122
                                                                                                6e-27
gi|625663|pir||JX0292 recA protein - Thermus aquaticus (strain HB8)
gi|1172880|sp|P42440|RECA_CAMJE RECA PROTEIN >gi|2119991|pir||I4...
                                                                                         121
                                                                                                le-26
                                                                                         120
                                                                                                2e-26
gi|4154654 (AE001453) RECA PROTEIN. [Helicobacter pylori J99]
gi|1072968|pir||C55020 recA protein - Thermus sp >gi|458472|dbj|...
                                                                                          120
                                                                                                2e-26
                                                                                         120
                                                                                                2e-26
gi|3219852|sp|P95469|RECA_PARDE RECA_PROTEIN >gi|1825468 (U59631...
gi|2507284|sp|P42445|RECA_HELPY_RECA_PROTEIN >gi|2313235|gb|AAD0...
                                                                                         119
                                                                                                3e-26
                                                                                                4e-26
                                                                                         119
gi|1172890 sp|Q02350 RECA_STAAU RECA PROTEIN >gi|463285 (L25893)...
                                                                                         118 Se-26
gi|4416209|gb|AAD20261| (AF094756) RecA protein (Bifidobacterium...
                                                                                                5e-26
                                                                                          118
gi|2500084|sp|Q59180|RECA_BORBU RECA PROTEIN >gi|1276443 (U23457...
                                                                                                5e-26
                                                                                          118
Query= sid|114832|lan|dp10RF011 Phage dp1 ORF|28017-29096|3
            (359 letters)
gi|2444110 (U88974) ORF31 (Streptococcus thermophilus temperate ...
                                                                                         187 le-46
gi|3320438 (AF057033) gp348 (Streptococcus thermophilus bacterio...
gi|479514|pir||S34244 hypothetical protein p38 - actinophage VWB...
                                                                                         179 2e-44
Query= sid|114834|lan|dplORF013 Phage dp1 ORF|10215-11240|3
            (341 letters)
gi|580855|emb|CAA29958| (X06803) dnaZX-like ORF put. DNA polymer...
                                                                                         182 2e-45
gi|118807|sp|P09122|DP3X_BACSU DNA POLYMERASE III SUBUNITS GAMMA...
                                                                                          182 2e-45
gi|98292|pir||S13786 DNA-directed DNA polymerase (EC 2.7.7.7) II...
                                                                                          182
                                                                                                2e-45
gi|1527142 (U66040) DNA polymerase III gamma subunit (Salmonella...
                                                                                                4e-42
                                                                                          172
gi 2494197 | sp | P74876 | DP3X_SALTY DNA POLYMERASE III SUBUNITS GAMM ...
                                                                                          172
                                                                                                4e-42
gi 118808 sp P06710 DP3X_ECOLI DNA POLYMERASE III SUBUNITS GAMMA...
                                                                                          170
                                                                                                1e-41
gi 4155207 (AE001497) DNA POLYMERASE III SUBUNITS GAMMA AND TAU ...
                                                                                          169
                                                                                                2e-41
gi 2313841 gb AAD07767.1 (AE000584) DNA polymerase III gamma an...
                                                                                          168
                                                                                                4e-41
gi 2583049 (AF025391) DNA polymerase III holoenzyme tau subunit ...
                                                                                         166
                                                                                                3e-40
gi|2884127 (AE000759) DNA polymerase III gamma subunit [Aquifex ... gi|3861390|emb|CAA15289| (AJ235273) DNA POLYMERASE III SUBUNITS ...
                                                                                         166
                                                                                                3e-40
                                                                                         165
                                                                                                5e-40
gi|1169397|sp|P43746|DP3X_HAEIN DNA POLYMERASE III SUBUNITS GAMM...
                                                                                          156
                                                                                                2e-37
gi|1293572 (U49738) DNA polymerase III tau homolog DnaX (Cauloba...
                                                                                          151
                                                                                                8e-36
gi|3328753 (AE001306) DNA Pol III Gamma and Tau (Chlamydia trach...
gi|4376294|gb|AAD18193| (AE001589) DNA Polymerase III Gamma and ...
gi|581255|emb|CAA28175| (X04487) alternate dnaZX protein (AA 1-6...
                                                                                          148
                                                                                                4e-35
                                                                                                5e-35
                                                                                          148
                                                                                                3e-34
                                                                                          146
gi|2688379 (AE001151) DNA polymerase III, subunits gamma and tau...
                                                                                                2e-32
                                                                                          140
gi|3323329 (AE001268) DNA polymerase III, subunits gamma and tau...
                                                                                          137
                                                                                                1e-31
Query= sid|114835|lan|dp10RF014 Phage dp1 ORF|50961-51974|3
            (337 letters)
gi|1346796|sp|P47492|PRIM_MYCGE DNA PRIMASE >gi|1361496|pir||F64...
                                                                                           57 2e-07
 gi|740008|prf||2004290A primase [Haemophilus influenzae]
                                                                                                le-05
gi|1172619|sp|Q08346|PRIM_HAEIN DNA PRIMASE >gi|1074033|pir||A64...
                                                                                                1e-05
gi 1709769 sp | Q04505 | PRIM_LACLA DNA PRIMASE >gi | 1075726 | pir | JC2...
                                                                                           51
                                                                                                1e-05
 gi|639846|dbj|BAA03516| (D14690) DNA primase [Lactococcus lactis]
                                                                                                le-05
Query= sid | 114837 | lan | dp1ORF016 Phage dp1 ORF | 43413-44303 | 3
            (296 letters)
 qi|1934766|emb|CAB07986| (Z93946) N-acetylmuramoyl-L-alanine ami...
                                                                                                0.0
 gi 113676 sp P06653 ALYS_STRPN AUTOLYSIN (N-ACETYLMURAMOYL-L-ALA...
                                                                                          221
                                                                                                46-57
gi 282326 pir A42935 N-acetylmuramoyl-L-alanine amidase (EC 3.5...
                                                                                          219
                                                                                                3e-56
 gi|416618|sp|P32762|ALYS_BPHB3 LYTIC AMIDASE (N-ACETYLMURAMOYL-L...
                                                                                          212
                                                                                                2e-54
gi|285273|pir||A42936 N-acetylmuramoyl-L-alanine amidase (EC 3.5...
                                                                                          212
                                                                                                26-54
 gi | 127787 | sp | P15057 | LYCA_BPCP1 LYSOZYME (ENDOLYSIN) (MURAMIDASE) ...
                                                                                          162
                                                                                                4e-39
gi|67761|pir||MUBPCP N-acetylmuramoyl-L-alanine amidase (EC 3.5....
                                                                                          162
                                                                                                4e-39
 gi|127789|sp|P19386|LYCA_BPCP9 LYSOZYME (ENDOLYSIN) (MURAMIDASE)...
                                                                                                1e-38
                                                                                          160
 gi|928832 (L44593) ORF259; putative [Lactococcus lactis phage BK...
                                                                                                2e-26
                                                                                          119
 gi|2511705|emb|CAA71783| (Y10818) sigA binding protein (Streptoc...
                                                                                                9e-24
                                                                                          111
 gi|4097980 (U72655) surface protein C [Streptococcus pneumoniae]
                                                                                                1e-22
                                                                                          107
 gi|2351768 (U89711) PapA (Streptococcus pneumoniae)
                                                                                          105
                                                                                                4e-22
 gi|2425109 (AF019904) choline binding protein A (Streptococcus p...
                                                                                                6e-22
                                                                                          104
 gi|282335|pir||A41971 surface protein pspA precursor - Streptoco...
                                                                                                1e-21
                                                                                          104
                                                                                                2e-21 ~
gi 2576331 emb CAA05158 (AJ002054) SpsA protein (Streptococcus ... gi 2127295 pir | S57962 cspC protein - Clostridium acetobutylicum...
                                                                                          103
                                                                                                6e-16
                                                                                           85
gi|2176333|emb|CAA05159| (AJ002055) SpsA protein (Streptococcus ... gi|4106522|gb|AAD02874.1| (AF097909) excreted protein FibB [Pept... gi|1361406|pir||557714 cspB protein - Clostridium acetobutylicum... gi|1914872|emb|CAB04758| (Z82001) PCPA [Streptococcus pneumoniae]
                                                                                                le-15
                                                                                           83 3e-15
                                                                                                4e-15
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gi|3168594|dbj|BAA28613| (AB012763) SpaA (Erysipelothrix rhusiop...
gi|2292750|emb|CAA64942| (X95646) homology to orf259 of lactococ...
                                                                                                   81 le-14
                                                                                                   80 3e-14
gi|2935696 (AF032122) putative lysin [Streptococcus thermophilus...
                                                                                                   80 3e-14
gi|4586910|dbj|BAA76540.1| (AB017447) protective antigen SpaA.1 ...
                                                                                                   80 3e-14
gi|3540294 (AF057033) lysin [Streptococcus thermophilus bacterio...
                                                                                                   79 5e-14
Query= sid|114841|1an|dp1ORF020 Phage dp1 ORF|1864-2658|1
             (264 letters)
gi|2633745|emb|CAB13247| (Z99111) similar to coenzyme PQQ synthe...
gi|2808502|emb|CAA12532| (AJ225561) ExsD protein [Sinorhizobium ...
gi|3861151|emb|CAA15051| (AJ235272) unknown [Rickettsia prowazekii]
gi|1652793|dbj|BAA17712| (D90908) hypothetical protein [Synechoc...
gi|1723815|sp|P55139|YGCF_ECOLI HYPOTHETICAL 25.0 KD PROTEIN IN ...
                                                                                                  217 5e-56
                                                                                                  163 le-39
                                                                                                   82 6e-15
                                                                                                   76 3e-13
                                                                                                   70
                                                                                                        2e-11
gi|2984272 (AE000769) hypothetical protein [Aquifex aeolicus]
gi|4155435 (AE001516) putative [Helicobacter pylori J99]
                                                                                                         4e-10
                                                                                                   66
                                                                                                   57
                                                                                                         le-07
gi|2127833|pir||C64505 coenzyme PQQ synthesis protein III homolo...
                                                                                                         5e-07
                                                                                                   55
gi|2622338 (AE000890) coenzyme PQQ synthesis protein III [Methan...
                                                                                                         9e-07
                                                                                                   54
gi|3257042|dbj|BAA29725| (AP000003) 254aa long hypothetical prot...
gi|2314068|gb|AAD07976.1| (AB000602) conserved hypothetical prot...
                                                                                                         2e-06
                                                                                                   53
                                                                                                         6e-06
                                                                                                   52
gi|1723816|sp|P45097|YGCF_HABIN HYPOTHETICAL PROTEIN HI1189 >gi|...
                                                                                                   50 2e-05
Query= sid|114842|lan|dp1ORF021 Phage dp1 ORF|2504-3295|2
gi|127481|sp|P19465|GCH1_BACSU GTP CYCLOHYDROLASE I (GTP-CH-I) >...
gi|3242315|emb|CAA04237| (AJ000685) GTP cyclohydrolase (Streptoc...
gi|2494695|sp|Q54769|GCH1_SYNP7 GTP CYCLOHYDROLASE I (GTP-CH-I) ...
                                                                                                  208 4e-53
                                                                                                         4e-48
gi|255061|bbs|112832 ($44049) GTP cyclohydrolase I (clone hGCH-1...
gi|4503949|ref|NP_000152.1|PGCH1| GTP cyclohydrolase I (dopa-res...
gi|2113967|emb|CAB08935| (295557) folE [Mycobacterium tuberculosis]
                                                                                                  187
 gi|1730240|sp|P50141|GCH1_CHICK GTP CYCLOHYDROLASE I (GTP-CH-I) ...
                                                                                                  185
 gi 2494696 sp Q55759 GCH1_SYNY3 GTP CYCLOHYDROLASE I (GTP-CH-I) ...
 gi|121061|sp|P22288|GCH1_RAT GTP CYCLOHYDROLASE I PRECURSOR (GTP...
 gi|3183014|sp|013774|GCH1_SCHPO GTP CYCLOHYDROLASE I (GTP-CH-I) ...
                                                                                                         6e-46
 gi|3097224|emb|CAA18795| (AL023093) GTP cyclohydrolase I [Mycoba...
                                                                                                  182 2e-45
 gi|2494697|sp|Q19980|GCH1_CAEEL PROBABLE GTP CYCLOHYDROLASE I (G...
                                                                                                         2e-45
                                                                                                  182
 gi | 462167 | sp | Q05915 | GCH1 MOUSE GTP CYCLOHYDROLASE I PRECURSOR (G... gi | 1669664 | emb | CAA89808 | (Z49706) GTP cyclohydrolase I [Dictyost...
                                                                                                  180 7e-45
                                                                                                  180 le-44
 gi|2981082 (AF052048) GTP-cyclohydrolase [Ostertagia ostertagi]
gi|31954|emb|CAA78908| (Z16418) GTP cyclohydrolase I [Homo sapi...
                                                                                                  178 3e-44
                                                                                                  177 8e-44
 gi|551344|bbs|150280 (S71373) GTP cyclohydrolase I [mice, Peptid...
gi|1730247|sp|P51601|GCH1_YEAST GTP CYCLOHYDROLASE I (GTP-CH-I) ...
                                                                                                  174 Se-43
                                                                                                  174
                                                                                                         7e-43
 gi|1246912|emb|CAA87397| (Z47201) GTP cyclohydrolase 1 [Saccharo...
gi|1730246|sp|P51595|GCH1_STRPN GTP CYCLOHYDROLASE I (GTP-CH-I) ...
                                                                                                  172 2e-42
                                                                                                  168
                                                                                                         3e-41
 gi|2982951 (AE000680) GTP cyclohydrolase I (Aquifex aeolicus)
                                                                                                        6e-40
                                                                                                  164
 Ouery= sid|114843|lan|dp10RF022 Phage dp1 ORF|30896-31675|2
              (259 letters)
 gi|2347102 (U77367) internalin [Listeria monocytogenes]
                                                                                                    55 5e-07
                                                                                                    52 4e-06
 gi|3123226|sp|P25146|INLA_LISMO INTERNALIN A PRECURSOR >gi|48705...
 gi 149674 (M67471) internalin [Listeria monocytogenes]
                                                                                                         4e-06
 Query= sid|114850|lan|dp10RF029 Phage dp1 ORF|662-1348|2
              (228 letters)
 gi|2650185 (AE001074) succinoglycan biosynthesis regulator (exsB...
 gi|3861231|emb|CAA15131| (AJ235272) unknown [Rickettsia prowazekii]
                                                                                                   117
                                                                                                         8e-26
 gi|2622210 (AE000881) conserved protein [Methanobacterium thermo...
                                                                                                   108 4e-23
 gi|2983380 (AE000709) trans-regulatory protein ExsB [Aquifex aeo...
gi|1001327|dbj|BAA10814| (D64006) ExsB (Synechocystis sp.)
                                                                                                    88 6e-17
                                                                                                         6e-17
 gi|2128055|pir||B64468 hypothetical protein homolog MJ1347 - Met...
gi|4155143 (AE001491) putative [Helicobacter pylori J99]
                                                                                                    83 le-15
                                                                                                         4e-15
 gi|2313760|gb|AAD07701.1| (AE000578) conserved hypothetical prot...
                                                                                                    80 2e-14
 gi|2120814|pir||S60183 protein ExsB - Rhizobium meliloti >gi|114...
                                                                                                    76
                                                                                                         3e-13
 gi|2633743|emb|CAB13245| (Z99111) similar to hypothetical protei...
gi|1175543|sp|P44124|YBAX_HAEIN HYPOTHETICAL PROTEIN HI1191 >gi|...
                                                                                                    75
                                                                                                         5e-13
                                                                                                    74 le-12
 gi|2495537|sp|P77756|YBAX_ECOLI HYPOTHETICAL 25.5 KD PROTEIN IN ...
gi|3256471|dbj|BAA29154.1| (AP000001) 269aa long hypothetical pr...
                                                                                                    67 1e-10 ---
                                                                                                    54 le-06
 gi|2921156 (AF022216) aluminum resistance protein (Arthrobacter ...
 Query= sid | 114855 | lan | dp1ORF034 Phage dp1 ORF | 131-652 | 2
              (173 letters)
                                                                                                  220 4e-57
 qi|2633746|emb|CAB13248| (Z99111) similar to hypothetical protei...
```

gi 4155926 (AE001554) putative [Helicobacter pylori J99] gi 2314588 gb AAD08456.1  (AE000642) conserved hypothetical prot gi 2983458 (AE000714) hypothetical protein [Aquifex aeolicus] gi 1006604 dbj BAA10757  (D64005) hypothetical protein [Synechoc gi 2967529 (U11045) unknown [Buchnera aphidicola] gi 2495654 sp Q46920 YQCD_ECOLI HYPOTHETICAL 32.6 KD PROTEIN IN gi 1175604 sp P44153 YQCD_HAEIN HYPOTHETICAL PROTEIN H11291 >gi  gi 3860642 emb CAA14543  (AJ235270) unknown [Rickettsia prowazekii] Query= sid 114857 lan dplORF036 Phage dpl ORF 48808-49362 1	162 161 103 87 79 69 63 56	1e-39 3e-39 9e-22 6e-17 2e-14 2e-11 1e-09 1e-07
(184 letters) gi 1353529 (U38906) ORF12 [Bacteriophage rlt] Query= sid 114859 lan dp1ORF038 Phage dp1 ORF 1350-1871 3 (173 letters)	53	le-06
gi 1175542 sp P44123 YB90_HAEIN HYPOTHETICAL PROTEIN HI1190 >gi  gi 2982977 (AE000681) hypothetical protein [Aquifex aeolicus] gi 3860744 emb CAA14645  (AJ235270) unknown [Rickettsia prowazekii] gi 2650193 (AE001074) conserved hypothetical protein [Archaeoglo gi 3258383 dbj BAA31066.1  (AP000007) 157aa long hypothetical pr gi 1001713 dbj BAA10550  (D64004) hypothetical protein [Synechoc gi 4155434 (AE001516) putative [Helicobacter pylori J99]	100 67 65 58 55 50	2e-07
Query= sid 114860 lan dp10RF039 Phage dp1 ORF 3306-3803 3 (165 letters)  gi 1922884 emb CAA68244  (X99978) ORF7; hydophobic protein [Lact  Query= sid 114862 lan dp10RF041 Phage dp1 ORF 8208-8699 3	64	5e-10
(163 letters)  gi 2522313 (AF012906) dUTPase homolog (Bacillus subtilis) >gi 26 gi 2634150 emb CAB13650  (Z99113) similar to deoxyuridine 5'-tri gi 3913546 sp 054134 DUT_STRCO DEOXYURIDINE 5'-TRIPHOSPHATE NUCL gi 3913542 sp 048500 DUT_BPTS DEOXYURIDINE 5'-TRIPHOSPHATE NUCLE gi 3913548 sp 068992 DUT_CHLTE DEOXYURIDINE 5'-TRIPHOSPHATE NUCL	108 108 56 52 50	3e-23
Query= sid 114867 lan dp10RF046 Phage dp1 ORF 42774-43202 3 (142 letters)  qi 1934764 emb CAB07984  (Z93946) hypothetical protein [bacterio	287	2e-77
Query= sid 114901 lan dp10RF080 Phage dp1 ORF 42490-42759 1 (89 letters)		
gi 1934763 emb CAB07983  (Z93946) hypothetical protein (bacterio  Query= sid 114912 lan dp10RF091 Phage dp1 ORF 43189-43413 1 (74 letters)	147	1e-35
gi 1934765 emb CAB07985  (293946) holin [bacteriophage Dp-1]	63	2e-10

- \_\_\_\_\_\_

Table 32

# Sequence of Dp1 published by Sheehan and al.. 4731 nucleotides.

•	•	•	_					
1						ccatgtattc		
71						ccttttagtg		
141	agaccttaaa	tatcgaattg	actcaaaagc	cgatcaaaag	ctaactaacc	aacagttgac	ggcactcacg	
211	gaaaaggctc	aactacatga	cgcagaactg	aaagctaagg	ctacaatgga	gcagttaagt	aacttagaaa	
281	aggcttatga	aggtagaatg	aaagctaatg	aagaagctat	caacaaatcg	gaacccgacc	taatcttagc	
351	ggcaagtcga	attgaagcta	ctatccaaga	acttggcggg	ctacgggaac	tgaagaagtt	cgtcgacagt	
421	tgcatgagct	cttctaatca	aggtctaatt	atcggtaaga	acgacggtag	ctctaccatt	aaggtatcaa	
491	gtgaccgaat	ttctatgttc	tccgcaggga	atgaagttat	gtaccttacg	caagggttca	ttcacatcga	
561						aatactcgtt		
631	atgaacgtga	ttcggtatgt	aggataagga	gaataacatg	acaaaattta	tcaactcata	cggccctctt	
701	cacttgaacc	tttacgtcga	acaagttagt	caggacgtaa	cgaacaactc	ctcgcgagtt	agttggcgag	
771	ctactgtcga	ccgcgatgga	gcttatcgaa	cgtggactta	tggaaatatt	agtaaccttt	ccgtatggtt	
841	aaatggttca	agtgttcata	gcagtcaccc	agactacgac	acgtccggcg	aagaggtaac	gctcgcaagt	
911	ggagaagtga	ctgttcctca	caatagtgac	gggacaaaga	caatgtccgt	ttgggcttcg	tttgacccta	
981	ataacggcgt	tcacggaaat	atcactatct	ctactaatta	cactttagac	agtattccaa	ggtctacaca	
1051	gatttctagt	tttgagggaa	atcgaaatct	aggatcttta	catacggtta	tctttaaccg	aaaagtgaac	
1121	tettttacge	atcaagtttg	gtaccgagtt	ttcggtagcg	actggataga	tttaggtaag	aaccatacta	
1191	ctagcgtatc	ctttacgccg	tcactggact	tagcaaggta	cttacctaaa	tcaagttccg	gaacaatgga	
1261	catctgtatt	cgaacctata	acggaactac	gcaaattggt	agtgacgtct	attcaaacgg	atggaggttc	
1331	aacatccccg	attcagtacg	tcctactttt	tcgggcattt	ctttagtaga	cacgacttca	gcggttcgac	
1401	agattttaac	agggaacaac	ttcctccaaa	tcatgtcgaa	cattcaagtc	aacttcaaca	atgcttccgg	
1471	cgcttacgga	tccactatcc	aagcatttca	cgctgagctc	gtaggtaaaa	accaagctat	caacgaaaac	
1541	ggcggcaaat	tgggtatgat	gaactttaat	ggctccgcta	ccgtaagagc	atgggttaca	gacacgcgag	
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1681						ctaaggtcgc		
1751						gttgaacact		
1821						actaactcgt		
1891						aagacaggtt		
1961						ggacggtcga		
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2451						atgtgcataa		
2521						ttcaatgtat		
2591						cttgtggtga		
2661						agctgaaatc		
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2801	attggacaag	gagctgaaaa	gaaacttgtc	aaaacaacga	ttgtgaacat	tgatgcaaac	gcagtatcaa	
2871	ccgtctctga	aactcttcat	gacccagact	tgtatgctgc	gaaccgtcga	gaacttcgag	ctgacgagca	
2941	aaaacttcgc	gaaactcgtt	acgcaatcga	agatgaaatt	aatagctgga	gcgggggaaa	aaagggggag	
3011	cccggctcta	acaggctgaa	taaggaggcg	tcaatctatg	ccaatgtggc	taaacgacac	cgcagtcttg	
3081	acgacgatta	ttacagcgtg	cagcggagtg	cttactgtcc	tactaaataa	gttattcgaa	tggaaatcga	
3151	ataaagccaa	gagcgtttta	gaggatatct	ctacaactct	tagcactctt	aaacagcagg	tcgacgggat	
3221	tgaccaaacg	acagtagcaa	tcaatcacca	aaatgacgtc	attcaagacg	gaactagaaa	aattcaacgt	
3291	taccgtcttt	atcacgactt	aaaaagggaa	gtgataacag	gctatacaac	tctcgaccat	tttagagagc	
3361	tctctatttt	attcgaaagt	tataagaacc	ttggcggaaa	tggtgaagtt	gaagccttgt	atgaaaaata	
3431	caagaaatta	ccaattaggg	aggaagattt	agatgaaact	atctaacgaa	caatatgacg	tagcaaagaa	
3501						gagcgttgta		
3571	actactgcta	tcacaggaac	cattgcactt	cttgcaactt	ttgcaggtac	tgttctagga	gtttctagcc	
3641	gaaactacca	aaaggaacaa	gaagctcaaa	acaatgaggt	ggaataatgg	gagtcgatat	tgaaaaaggc	
3711	gttgcgtgga	tgcaggcccg	aaagggtcga	gtatcttata	gcatggactt	tcgagacggt	cctgatagct	Ĺ
3781						agtgctggat		`
3851						gtgaaaatgc		
3921						cgctggaggt		
3991						atttccgtca		
4061						ctaacgcaaa		
4131						tcgagcaaac		
4201						gacgaccaag		
4271	cgctgagaaa	tggttgaaac	atactgatgg	aaattggtat	tggttcgacc	gtgacggata	catggctacg	 
4341							ggttggatta-	
4411						tcgaatgcgt		
4481						ctcaattcac		
4551						ttcttaatat		
4621						atttacttat	tcgaagattt	
4691	caattataat	taaataatca	acgagattca	taattggagg	aatg			

### Table 33

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gi|4538725|emb|AJ240754.1|SPN240754 gi|4519233|dbj|AB011207.1|AB011207 [4519233] [4538725] gi|4519231|dbj|AB011206.1|AB011206 [4519231] gi|4538722|emb|AJ240753.1|SPN240753 gi|4519229|dbj|AB011205.1|AB011205 [4519229] [4538722] gi|4519227|dbi|AB011204.1|AB011204 [4519227] gi|4538719|emb|AJ240752.1|SPN240752 [4538719] gi|4519225|dbj|AB011203.1|AB011203 [4519225] gi|4538716|emb|AJ240751.1|SPN240751 gi|4519223|dbj|AB011202.1|AB011202 [4519223] [4538716] gi|4519221|dbj|AB011201.1|AB011201 [4519221] gi|4538713|emb|AJ240750.1|SPN240750 gi|4519219|dbj|AB011200.1|AB011200 [4519219] [4538713] gi|4519217|dbj|AB011199.1|AB011199 [4519217] gi|4538710|emb|AJ240749.1|SPN240749 [4538710] gi|4519215|dbj|AB011198.1|AB011198 [4519215] gi|4538707|emb|AJ240748.1|SPN240748 gi|4495127|emb|AJ240605.1|SPN240605 [4538707] [4495127] gi|4538704|emb|AJ240747.1|SPN240747 gi|4468031|emb|AJ132957.1|SPN132957 [4538704] [4468031] gi|4538701|emb|AJ240746.1|SPN240746 gi|4468029|emb|AJ132956.1|SPN132956 [4538701] [4468029] gi|4538698|emb|AJ240745.1|SPN240745 gi|4218532|emb|AJ010312.1|SPN010312 [4538698] [4218532] gi|4538695|emb|AJ240744.1|SPN240744 gi|4456852|emb|AJ236792.1|SPN236792 [4538695] [4456852] gi|4538692|emb|AJ240743.1|SPN240743 gi|4456850|emb|AJ236791.1|SPN236791 [4538692] [4456850] gi|4538689|emb|AJ240742.1|SPN240742 gi|4456848|emb|AJ236790.1|SPN236790 [4538689] [4456848] gi|4538686|emb|AJ240741.1|SPN240741 gi|4456846|emb|AJ236789.1|SPN236789 [4538686] [4456846] gi|4538683|emb|AJ240740.1|SPN240740 gi|3550644|emb|AJ006987.1|SPAJ6987 [3550644] [4538683] gi|3550625|emb|AJ006986.1|SPAJ6986 [3550625] gi|4538680|emb|AJ240739.1|SPN240739 gi|4416518|gb|AF014458.2|AF014458 [4416518] [4538680] gi|4406260|gb|AF105116.1|AF105116 [4406260] gi|4538677|emb|AJ240738.1|SPN240738 [4538677] gi|4406257|gb|AF105115.1|AF105115 [4406257] gi|4530444|gb|AF118229.1|AF118229 [4530444] gi|4406254|gb|AF105114.1|AF105114 [4406254] gi|4519253|dbj|AB015852.1|AB015852 [4519253] gi|4406246|gb|AF105113.1|AF105113 [4406246] gi|4519251|dbj|AB015851.1|AB015851 [4519251] gi|4406243|gb|AF105112.1|AF105112 [4406243] gi|4519249|dbj|AB015850.1|AB015850 [4519249] gi|4138533|emb|AJ005815.1|SPN5815 [4138533] gi|4519247|dbj|AB015849.1|AB015849 [4519247] gij3821726|emb|AJ232433.1|SPN232433 [3821726] gi|4519245|dbj|AB015848.1|AB015848 [4519245] gi|3821724|emb|AJ232432.1|SPN232432 gi|4519243|dbj|AB015847.1|AB015847 [4519243] [3821724] gi|4519241|dbj|AB015846.1|AB015846 [4519241] gi|3821722|emb|AJ232431.1|SPN232431gi|4519239|dbj|AB011210.1|AB011210 [4519239] [3821722] gi|4519237|dbj|AB011209.1|AB011209 [4519237] gi|3821720|emb|AJ232430.1|SPN232430 [3821720] gi|4519235|dbj|AB011208.1|AB011208 [4519235]

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gi 3821424 emb AJ232276.1 SPN232276 [3821424]		gi 3821376 emb AJ232252.1 SPN232252 [3821376]
gi 3821422 emb AJ232275.1 SPN232275 [3821422]		gi 3821374 emb AJ232251.1 SPN232251 [3821374]
gi 3821420 emb AJ232274.1 SPN232274 [3821420]		gi 3821372 emb AJ232250.1 SPN232250 [3821372]
gi 3821418 emb AJ232273.1 SPN232273 [3821418]		gi 3821370 emb AJ232249.1 SPN232249 [3821370]
gi 3821416 emb AJ232272.1 SPN232272 [3821416]		gi 3821367 emb AJ232248.1 SPN232248 [3821367]
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gi 3821408 emb AJ232268.1 SPN232268 [3821408]		gij3821359 emb AJ232244.1 SPN232244 [3821359]
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gi 3821404 emb AJ232266.1 SPN232266 [3821404]		gi 3821355 emb AJ232241.1 SPN232241 [3821355]
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[3821402]		gi 2909863 gb AF047696.1 AF047696 [2909863]
gi 3821400 emb AJ232264.1 SPN232264 [3821400]		gi 4193353 gb AF055088.1 AF055088 [4193353]
gi 3821398 emb AJ232263.1 SPN232263 [3821398]		gi 4185242 gb AH007276.1 SEG_SPTNJUNC [4185242]
gi 3821396 emb AJ232262.1 SPN232262 [3821396]		gi 4185241 gb AF066797.1 SPTNJUNC2 [4185241]
gi 3821394 emb AJ232261.1 SPN232261 [3821394]		gi 4185240 gb AF066796.1 SPTNJUNC1 [4185240]
gi 3821392 emb AJ232260.1 SPN232260 [3821392]		gi 4097979 gb U72655.1 SPU72655 [4097979] gi 4063720 gb L29323.1 STRMTR [4063720]
gi 3821390 emb AJ232259.1 SPN232259 [3821390]		gi 1657605 gb U66846.1 SPU66846 [1657605] gi 1657602 gb U66845.1 SPU66845_[1657602]
gi 3821388 emb AJ232258.1 SPN232258 [3821388]		gi 4009485 gb AF068903.1 AF068903 [4009485]
gi 3821386 emb AJ232257.1 SPN232257 [3821386]		gi 4009477 gb AF068902.1 AF068902 [4009477] gi 4009462 gb AF068901.1 AF068901 [4009462]

gi]3947767|emb|AJ233896.1|SPN233896 [3947767] gi|3947765|emb|AJ233895.1|SPN233895 [3947765] gi|3947763|emb|AJ233894.1|SPN233894 [3947763] gi|3947761|emb|AJ233893.1|SPN233893 [3947761] gi|3947759|emb|AJ233892.1|SPN233892 [3947759] gi|3947757|emb|AJ233891.1|SPN233891 [3947757] gi|3947755|emb|AJ233890.1|SPN233890 [3947755] gi|3947753|emb|AJ233889.1|SPN233889 [3947753] gi|3947751|emb|AJ233888.1|SPN233888 [3947751] gi|3947749|emb|AJ233887.1|SPN233887 [3947749] gi|3947730|emb|AJ233886.1|SPN233886 [3947730] gi|3758891|emb|Z71552.1|SPADCA [3758891] gi|3818479|gb|AF057294.1|AF057294 [3818479] gi|2351767|gb|U89711.1|SPU89711 [2351767] gi|3395661|dbj|AB006879.1|AB006879 [3395661] gi|3395659|dbj|AB006878.1|AB006878 [3395659] gi|3395657|dbj|AB006877.1|AB006877 [3395657] gi|3395655|dbj|AB006876.1|AB006876 [3395655] gi|3395653|dbj|AB006875.1|AB006875 [3395653] gi|3395651|dbj|AB006874.1|AB006874 [3395651] gi|3395649|dbj|AB006873.1|AB006873 [3395649] gi|3395647|dbj|AB006872.1|AB006872 [3395647] gi|3395645|dbj|AB006871.1|AB006871 [3395645] gi|3395643|dbj|AB006870.1|AB006870 [3395643] gi|3395641|dbj|AB006869.1|AB006869 [3395641] gi|3395639|dbj|AB006868.1|AB006868 [3395639] gi|2315992|gb|U87092.1|SPU87092 [2315992] gi|2209338|gb|U93576.1|SPU93576 [2209338] gi|2109442|gb|AF000658.1|SPDNAARG [2109442] gi|1881538|gb|U09239.1|SPU09239 [1881538] gi|1666904|gb|U76218.1|SPU76218 [1666904] gi|1613766|gb|U33315.1|SPU33315 [1613766]

gi|1498294|gb|U41735.1|SPU41735 [1498294] gi|1213493|gb|U47687.1|SPU47687 [1213493] gi|1163109|gb|U43526.1|SPU43526 [1163109] gi|556001|gb|U15171.1|SPU15171 [556001] gi|455063|gb|U02920.1|SPU02920 [455063] gi|784896|gb|L36923.1|STRSTRH [784896] gi|3320386|gb|AF030373.1|AF030373 [3320386] gi|2804772|gb|AF030374.1|AF030374 [2804772] gi|2804762|gb|AF030372.1|AF030372 [2804762] gi|2804756|gb|AF030371.1|AF030371 [2804756] gi|2804750|gb|AF030370.1|AF030370 [2804750] gi|2804745|gb|AF030369.1|AF030369 [2804745] gi|2804739|gb|AF030368.1|AF030368 [2804739] gi|2804732|gb|AF030367.1|AF030367 [2804732] gi|2804726|gb|AF030366.1|AF030366 [2804726] gi|2804720|gb|AF030365.1|AF030365 [2804720] gi|2804713|gb|AF030364.1|AF030364 [2804713] gi|2804707|gb|AF030363.1|AF030363 [2804707] gi|2804701|gb|AF030362.1|AF030362 [2804701] gi|2804694|gb|AF030361.1|AF030361 [2804694] gi|2804688|gb|AF030360.1|AF030360 [2804688] gi|2804682|gb|AF030359.1|AF030359 [2804682] gi|3550979|dbj|AB010387.1|AB010387 [3550979] gi|2275100|emb|AJ000336.1|SPR6LDH [2275100] gi|3551853|gb|AF076029.1|AF076029 [3551853] gi|3551773|gb|U94770.1|SPU94770 [3551773] gi|3550617|emb|AJ004869.1|SPAJ4869 [3550617] gi|3513563|gb|AF055727.1|AF055727 [3513563] gi|3513561|gb|AF055726.1|AF055726 [3513561] gi|3513559|gb|AF055725.1|AF055725 [3513559] gi|3513557|gb|AF055724.1|AF055724 [3513557] gi|3513555|gb|AF055723.1|AF055723 [3513555] gi|3513553|gb|AF055722.1|AF055722 [3513553] gi|3513549|gb|AF055721.1|AF055721 [3513549] gi|3513545|gb|AF055720.1|AF055720 [3513545] gi|1914869|emb|Z82001.1|SPZ82001 [-1914869] gi|2911421|gb|AF046238.1|AF046238 [2911421] gi|2911419|gb|AF046237.1|AF046237 [2911419] gi|2911417|gb|AF046236.1|AF046236 [2911417] gi|2911415|gb|AF046235.1|AF046235 [2911415]

gi|2911413|gb|AF046234.1|AF046234 [2911413] gi|2911411|gb|AF046233.1|AF046233 [2911411] gi|2911409|gb|AF046232.1|AF046232 [2911409] gi|2911407|gb|AF046231.1|AF046231 [2911407] gi|2911405|gb|AF046230.1|AF046230 [2911405] gi|3258601|gb|U40786.1|SPU40786 [3258601] gi|3211756|gb|AF052209.1|AF052209 [3211756] gi|3211752|gb|AF052208.1|AF052208 [3211752] gi|3211747|gb|AF052207.1|AF052207 [3211747] gi|3220194|gb|AF053121.1|AF053121 [3220194] gi|2766052|emb|Z99863.1|SPZ99863~[2766052]gi|2766050|emb|Z99862.1|SPZ99862 [2766050] gi|2766048|emb|Z99861.1|SPZ99861 [2766048] gi|2766046|emb|Z99860.1|SPZ99860 [2766046] gi|2766044|emb|Z99859.1|SPZ99859 [2766044] gi|2766042|emb|Z99858.1|SPZ99858 [2766042] gi|2766040|emb|Z99857.1|SPZ99857 [2766040] gi|2766038|emb|Z99856.1|SPZ99856 [2766038] gi|2766036|emb|Z99855.1|SPZ99855 [2766036] gi|2766034|emb|Z99854.1|SPZ99854 [2766034] gi|2766032|emb|Z99853.1|SPZ99853 [2766032] gi|2766030|emb|Z99852.1|SPZ99852 [2766030] gi|2766028|emb|Z99851.1|SPZ99851 [2766028] gi|2766026|emb|Z99850.1|SPZ99850 [2766026] gi|2766024|emb|Z99849.1|SPZ99849 [2766024] gi|2766022|emb|Z99848.1|SPZ99848 [2766022] gi|2766020|emb|Z99847.1|SPZ99847 [2766020] gi|2766018|emb|Z99846.1|SPZ99846 [2766018] gi|2766016|emb|Z99845.1|SPZ99845 [2766016] gi|2766014|emb|Z99844.1|SPZ99844 [2766014] gi|2766012|emb|Z99843.1|SPZ99843 [2766012] gi|2766010|emb|Z99842.1|SPZ99842 [2766010] gi|2766008|emb|Z99841.1|SPZ99841 [2766008] gi|2766006|emb|Z99840.1|SPZ99840 [2766006] gi|2766004|emb|Z99839.1|SPZ99839 [2766004] gi|2766002|emb|Z99838.1|SPZ99838 [2766002] gi|2766000|emb|Z99837.1|SPZ99837 [2766000] gi|2765998|emb|Z99828.1|SPZ99828 [2765998] gi|2765996|emb|Z99827.1|SPZ99827 [2765996] gi|2765994|emb|Z99826.1|SPZ99826 [2765994]

gi|2765992|emb|Z99825.1|SPZ99825 [2765992] gi|2765990|emb|Z99824.1|SPZ99824 [2765990] gi|2765988|emb|Z99823.1|SPZ99823 [2765988] gi|2765986|emb|Z99822.1|SPZ99822 [2765986] gi|2765984|emb|Z99821.1|SPZ99821 [2765984] gi|2765982|emb|Z99820.1|SPZ99820 [2765982] gi|2765980|emb|Z99819.1|SPZ99819 [2765980] gi|2765978|emb|Z99818.1|SPZ99818 [2765978] gi|2765976|emb|Z99817.1|SPZ99817 [2765976] gi|2765974|emb|Z99816.1|SPZ99816 [2765974] gi]2765972|emb|Z99815.1|SPZ99815 [2765972] gi|2765970|emb|Z99814.1|SPZ99814 [2765970] gi|2765968|emb|Z99813.1|SPZ99813 [2765968] gi|2765966|emb|Z99812.1|SPZ99812 [2765966] gi|2765964|emb|Z99811.1|SPZ99811 [2765964] gi|2765962|emb|Z99810.1|SPZ99810 [2765962] gi|2765960|emb|Z99809.1|SPZ99809 [2765960] gi|2765958|emb|Z99808.1|SPZ99808 [2765958] gi|2765956|emb|Z99807.1|SPZ99807 [2765956] gi|2765954|emb|Z99806.1|SPZ99806 [2765954] gi|2765952|emb|Z99805.1|SPZ99805 [2765952] gi|2765950|emb|Z99804.1|SPZ99804 [2765950] gi|2765948|emb|Z99803.1|SPZ99803 [2765948] gi|2894104|emb|X77249.1|SPR6CIARH [2894104] gi|3153897|gb|AF067128.1|AF067128 [3153897] gi|3152712|gb|AF065153.1|AF065153 [3152712] gi|3152710|gb|AF065152.1|AF065152 [3152710] gi|3152708|gb|AF065151.1|AF065151 [3152708] gi|3116426|gb|U84387.1|SPU84387 [3116426] gi|2385403|emb|AJ001247.1|SP7465RR3 [2385403] gi|2342540|emb|AJ001250.1|SP7978RR5 [2342540] gi|2342539|emb|AJ001251.1|SP7978RR3 [2342539] gi|2342538|emb|AJ001248.1|SP7466RR5 [2342538] gi|2342537|emb|AJ001249.1|SP7466RR3 [2342537] gi|3065896|gb|AF058920.1|AF058920 [3065896] gi|2982647|emb|AJ002294.1|SPAJ2294 [2982647]

gi|2982645|emb|AJ002293.1|SPAJ2293 [2982645] gi|2982643|emb|AJ002292.1|SPAJ2292 [2982643] gi|2982641|emb|AJ002291.1|SPAJ2291 [2982641] gi|1620466|emb|X99400.1|SPDACAO [1620466] gi|2196665|emb|Z84381.1|HSZ84381 [2196665] gi|2196663|emb|Z84380.1|HSZ84380 [2196663] gi|2196661|emb|Z84379.1|HSZ84379 [2196661] gi|2196659|emb|Z84378.1|HSZ84378 [2196659] gi|625175|gb|L36131.1|STREXP10A [625175] gi|3004945|gb|AF036624.1|AF036624 [3004945] gi|3004943|gb|AF036623.1|AF036623 [3004943] gi|3004941|gb|AF036622.1|AF036622 [3004941] gi|3004939|gb|AF036621.1|AF036621 [3004939] gi|3004937|gb|AF036620.1|AF036620 [3004937] gi|3004935|gb|AF036619.1|AF036619 [3004935] gij2370572|emb|Z86112.1|SPZ86112 [2370572] gi|2765946|emb|Z99802.1|SPZ99802 [2765946] gi|2398824|emb|Z34303.1|SPCINREC [2398824] gi|2894512|emb|AJ223491.1|SPPPR3 [2894512] gi|2198539|emb|X85787.1|SPCPS14E [2198539] gi|2766156|emb|Z99915.1|SPZ99915 [2766156] gi|2766154|emb|Z99914.1|SPZ99914 [2766154] gi|2766152|emb|Z99913.1|SPZ99913 [2766152] gi|2766150|emb|Z99912.1|SPZ99912 [2766150] gi|2766148|emb|Z99911.1|SPZ99911 [2766148] gi|2766146|emb|Z99910.1|SPZ99910 [2766146] gi|2766144|emb|Z99909.1|SPZ99909 [2766144] gi|2766142|emb|Z99908.1|SPZ99908 [2766142] gi|2766140|emb|Z99907.1|SPZ99907 [2766140] gi|2766138|emb|Z99906.1|SPZ99906 [2766138] gi|2766136|emb|Z99905.1|SPZ99905 [2766136] gi|2766134|emb|Z99904.1|SPZ99904 [2766134] gi|2766132|emb|Z99903.1|SPZ99903 [2766132] gi|2766130|emb|Z99902.1|SPZ99902 [2766130] gi|2766128|emb|Z99901.1|SPZ99901 [2766128] gi|2766126|emb|Z99900.1|SPZ99900 [2766126] gi|2766124|emb|Z99899.1|SPZ99899 [2766124] gi|2766122|emb|Z99898.1|SPZ99898 [2766122] gi|2766120|emb|Z99897.1|SPZ99897 [2766120] gi|2766118|emb|Z99896.1|SPZ99896 [2766118]

gi|2766116|emb|Z99895.1|SPZ99895 [2766116] gi|2766114|emb|Z99894.1|SPZ99894 [2766114] gi|2766112|emb|Z99893.1|SPZ99893 [2766112] gi|2766110|emb|Z99892.1|SPZ99892 [2766110] gi|2766108|emb|Z99891.1|SPZ99891 [2766108] gi|2766106|emb|Z99890.1|SPZ99890 [2766106] gi|2766104|emb|Z99889.1|SPZ99889 [2766104] gi|2766102|emb|Z99888.1|SPZ99888 [2766102] gi|2766100|emb|Z99887.1|SPZ99887 [2766100] gi|2766098|emb|Z99886.1|SPZ99886 [2766098] gi|2766096|emb|Z99885.1|SPZ99885 [2766096] gi|2766094|emb|Z99884.1|SPZ99884 [2766094] gi|2766092|emb|Z99883.1|SPZ99883 [2766092] gi|2766090|emb|Z99882.1|SPZ99882 [2766090] gi|2766088|emb|Z99881.1|SPZ99881 [2766088] gi|2766086|emb|Z99880.1|SPZ99880 [2766086] gi|2766084|emb|Z99879.1|SPZ99879 [2766084] gi|2766082|emb|Z99878.1|SPZ99878 [2766082] gi|2766080|emb|Z99877.1|SPZ99877 [2766080] gi|2766078|emb|Z99876.1|SPZ99876 [2766078] gi|2766076|emb|Z99875.1|SPZ99875 [2766076] gi|2766074|emb|Z99874.1|SPZ99874 [2766074] gi|2766072|emb|Z99873.1|SPZ99873 [2766072] gi|2766070|emb|Z99872.1|SPZ99872 [2766070] gi|2766068|emb|Z99871.1|SPZ99871 [2766068] gi|2766066|emb|Z99870.1|SPZ99870 [2766066] gi|2766064|emb|Z99869.1|SPZ99869 [2766064] gi|2766062|emb|Z99868.1|SPZ99868 [2766062] gi|2766060|emb|Z99867.1|SPZ99867 [2766060] gi|2766058|emb|Z99866.1|SPZ99866 [2766058] gi|2766056|emb|Z99865.1|SPZ99865 [2766056] gi|2766054|emb|Z99864.1|SPZ99864 [2766054] gi|2765906|emb|Z99206.1|SPZ99206 [2765906] gi|2765904|emb|Z99205.1|SPZ99205 [2765904] gi|2765902|emb|Z99204.1|SPZ99204 [2765902] gi|2765900|emb|Z99203.1|SPZ99203 [2765900] " gi|2765898|emb|Z99202.1|SPZ99202 [2765898] gi|2765896|emb|Z99201.1|SPZ99201 [2765896] gi|2765894|emb|Z99200.1|SPZ99200 [2765894] gi|2708631|gb|AF036951.1|AF036951 [2708631]

gi|886956|emb|Z49097.1|SPCS1112X [886956] gi|2656093|gb|L21856.1|STRMALR [2656093] gi|2576332|emb|AJ002055.1|SPSPSA47 [2576332] gi|2576330|emb|AJ002054.1|SPSPSA2 [2576330] gi|2511704|emb|Y10818.1|SPY10818 [2511704] gi|1944619|emb|Z83335.1|SPZ83335 [1944619] gi|2425108|gb|AF019904.1|AF019904 [2425108] gi|2385404|emb|AJ001246.1|SP7465RR5 [2385404] gi|438213|emb|Z16082.1|PNALIB [438213] gi|2149613|gb|U90721.1|SPU90721 [2149613] gi|49391|emb|Z21841.1|SPPBP2BB [49391] gi|2209207|gb|AF004325.1|AF004325 [2209207] gi|2293061|emb|Z95914.1|SPZ95914 [2293061] gi|2276393|gb|U16156.1|SPU16156 [2276393] gi|2183314|gb|AF003930.1|AF003930 [2183314] gi|2182093|emb|X95717.1|SPPARECGN [2182093] gi|984230|emb|Z49095.1|SPCS1111A [984230] gi|886954|emb|Z49096.1|SPCS1092X [886954] gi|1181613|dbj|D82873.1|STRPBP2BE [1181613] gi|1181612|dbj|D82871.1|STRPBP2BCZ [1181612] gi|1181611|dbj|D82870.1|STRPBP2BB2 [1181611] gi|1181579|dbj|D82869.1|STRPBP2BA1 [1181579] gi|1181192|dbj|D82872.1|STRPBP2BD [1181192] gi|575595|dbj|D42075.1|STRPBP2B2 [575595] gi|1339971|dbj|D42074.1|STRPBP2B1 [1339971] gi|2108329|emb|Y11463.1|SPDNAGCPO [2108329] gi|1944115|dbj|AB002522.1|AB002522 [1944115] gi|1666669|emb|Z77727.1|SPIS1381C [1666669] gi|1666668|emb|Z77726.1|SPIS1381B [1666668] gi|1666667|emb|Z77725.1|SPIS1381A [1666667] gi|1914873|emb|Z82002.1|SPZ82002 [1914873] gi|1431584|emb|Z74778.1|SPDHFR [1431584] gi|47452|emb|Z15120.1|SPSTRG [47452] gi|581717|emb|Z12159.1|SPCP131G [581717] gi|47342|emb|X17337.1|SPAMILOC [47342] gi|1800300|gb|U83667.1|SPU83667 [1800300] gi|1532066|emb|Y07780.1|SPTET0GEN [1532066]

gi|1460093|emb|X94909.1|SPIGA1PRT [1460093] gi|1750263|gb|U72720.1|SPU72720 [1750263] gi|298649|gb|S56948.1|S56948 [298649] gi|254537|gb|S43511.1|S43511 [254537] gi|245227|gb|S81051.1|S81051 [245227] gi|245226|gb|S81045.1|S81045 [245226] gi|245225|gb|S81043.1|S81043 [245225] gi|1150618|emb|Z49988.1|SPMMSAGEN [1150618] gi|47456|emb|X01138.1|SPTN917A [47456] gi|1658316|emb|Z47210.1|SPDEXCAP [1658316] gi|1550802|emb|X95385.1|SPCOMCGEN [1550802] gi|47457|emb|X01137.1|SPTN917B [47457] gi|975714|emb|X90941.1|SPTRJ5251 [975714] gi|975713|emb|X90940.1|SPTLJ5251 [975713] gi|975709|emb|X90939.1|SPDNATETM [975709] gi|1524346|emb|Z79691.1|SOORFS [1524346] gi|1553054|emb|X98364.1|SPPBPHU9 [1553054] gi|1553052|emb|X98367.1|SPPBPHU13 [1553052] gi|1553050|emb|X98366.1|SPPBPHU12 [1553050] gi|1553048|emb|X98365.1|SPPBPHU11 [1553048] gi|1575029|gb|U53509.1|SPU53509 [1575029] gi|1542968|gb|U49088.1|SPU49088 [1542968] gi|1542966|gb|U49087.1|SPU49087 [1542966] gi|1536961|emb|Y07845.1|SPGYRA [1536961] gi|47391|emb|X16367.1|SPPBPX [47391] gi|1490398|emb|Z67739.1|SPPARCETP [1490398] gi|1490395|emb|Z67740.1|SPGYRBORF [1490395] gi|1431589|emb|Z74777.1|SPTMRDHFR [1431589] gi|408145|emb|Z21702.1|SPUNGMUTX [408145] gi|47461|emb|X61025.1|SPXISINT [47461] gi|47459|emb|X55651.1|SPUNGG [47459] gi|47454|emb|X52632.1|SPT1545E [47454] gi|47421|emb|Z17307.1|SPRECA [47421] gi|47419|emb|X67873.1|SPPONA8 [47419] gi|47417|emb|X67872.1|SPPONA7 [47417] gi|47415|emb|X67871.1|SPPONA6 [47415]

gi|1161269|gb|L39074.1|STRSPXB [1161269]

gi|47413|emb|X67870.1|SPPONA5 [47413] gi|47411|emb|X67869.1|SPPONA4 [47411] gi|47409|emb|X67867.1|SPPONA2 [47409] gi|47407|emb|X67866.1|SPPONA1 [47407] gi|47405|emb|X67868.1|SPPNA3 [47405] gi|47403|emb|X52474.1|SPPLY [47403] gi|984232|emb|X16022.1|SPPENA [984232] gi|517190|emb|X78215.1|SPPBPXG [517190] gi|295840|emb|Z22230.1|SPPBP2BBA [295840] gi|288981|emb|Z22185.1|SPPBP2BAC [288981] gi|288979|emb|Z22184.1|SPPBP2BAB [288979] gi|288466|emb|Z21981.1|SPPBP2BAA [288466] gi|49390|emb|Z21813.1|SPPBP2XD [49390] gi|49389|emb|Z21812.1|SPPBP2XC [49389] gi|49387|emb|Z21811.1|SPPBP2BJ [49387] gi|49385|emb|Z21810.1|SPPBP2BI [49385] gi|49382|emb|Z21808.1|SPPBP2BH [49382] gi|49380|emb|Z21807.1|SPPBP2BG [49380] gi|49379|emb|Z21806.1|SPPBP2BF [49379] gi|49377|emb|Z21805.1|SPPBP2BE [49377] gi|49376|emb|Z21804.1|SPPBP2XB [49376] gi|49375|emb|Z21803.1|SPPBP2XA [49375] gi|49374|emb|Z21802.1|SPPBP2BD [49374] gi|49372|emb|Z21801.1|SPPBP2BC [49372] gi|49369|emb|Z21799.1|SPPBP2BA [49369] gi|47399|emb|X13137.1|SPPENASE [47399] gi|47397|emb|X13136.1|SPPENARE [47397] gi|1052802|emb|X83917.1|SPGYRBG [1052802] gi|587550|emb|X72967.1|SPNANA [587550] gi|49384|emb|Z21809.1|SPPBP1AB [49384] gi|49371|emb|Z21800.1|SPPBP1AA [49371] gi|984228|emb|Z49094.1|SPCS1091A [984228] gi|47372|emb|X54225.1|SPENDA [47372] gi|806590|emb|Z49246.1|SP667SOD [806590] gi|407172|emb|Z26851.1|SPATPAS2 [407172] gi|407166|emb|Z26850.1|SPATPAS1 [407166] gi|47353|emb|X63602.1|SPBOX [47353] gi|47348|emb|X05577.1|SPAPHA3 [47348] gi|47337|emb|X65132.1|SP824PBPX [47337] gi|47335|emb|X65134.1|SP669PBPX [47335]

gi|47331|emb|X65133.1|SP577PBPX [47331] gi|559527|emb|X65136.1|SP110PBPX [559527] gi|311415|emb|Z22807.1|SP16SRNAA [311415] gi|47329|emb|X65135.1|SP531PBPX [47329] gi|47307|emb|X65131.1|SP290PBPX [47307] gi|47295|emb|X58312.1|SP16SRNA [47295] gi|854614|emb|Z49109.1|SPGADAGN [854614] gi|556428|gb|L36660.1|STRORF1 [556428] gi|511062|emb|Z35135.1|SPALIAG [511062] gi|1208737|gb|U47625.1|SPU47625 [1208737] gi|530062|gb|U12567.1|SPU12567 [530062] gi|153656|gb|M29686.1|STRHEXB [153656] gi|153654|gb|M18729.1|STRHEXA [153654] gi|153608|gb|M14339.1|STRDPN2A [153608] gi|153605|gb|M14340.1|STRDPN1A [153605] gi|643543|gb|U20084.1|SPU20084 [643543] gi|643541|gb|U20083.1|SPU20083 [643541] gi|643539|gb|U20082.1|SPU20082 [643539] gi|643537|gb|U20081.1|SPU20081 [643537] gi|643535|gb|U20080.1|SPU20080 [643535] gi|643533|gb|U20079.1|SPU20079 [643533] gi|643531|gb|U20078.1|SPU20078 [643531] gi|643529|gb|U20077.1|SPU20077 [643529] gi|643527|gb|U20076.1|SPU20076 [643527] gi|643525|gb|U20075.1|SPU20075 [643525] gi|643523|gb|U20074.1|SPU20074 [643523] gi|643521|gb|U20073.1|SPU20073 [643521] gi|643519|gb|U20072.1|SPU20072 [643519] gi|643517|gb|U20071.1|SPU20071 [643517] gi|643515|gb|U20070.1|SPU20070 [643515] gi|643513|gb|U20069.1|SPU20069 [643513] gi|643511|gb|U20068.1|SPU20068 [643511] gi|643509|gb|U20067.1|SPU20067 [643509] gi|1017802|gb|U37560.1|SPU37560 [1017802] gi|663277|gb|M36180.1|STRCOMAA [663277] gi|437704|gb|L20670.1|STRHYALURO [437704] gi|153849|gb|L07751.1|TRNTN5252R [153849] gi|153855|gb|M25519.1|STRVA1 [153855] gi|153853|gb|M80215.1|STRUVS402A [153853] gi|153848|gb|L07750.1|STRTN5252L [153848]

gi]153840|gb|M74122.1|STRSURPROA [153840] gi|153796|gb|M60763.1|STRRRNAA [153796] gi|153791|gb|M31296.1|STRRECP [153791] gi|516639|gb|L20556.1|STRPLPA [516639] gi|153783|gb|M28679.1|STRPROMB [153783] gi|153782|gb|M28678.1|STRPROMA [153782] gi|153766|gb|M90527.1|STRPONA [153766] gi|153764|gb|J04479.1|STRPOLA [153764] gi|153752|gb|M25515.1|STRNG4369 [153752] gi|153722|gb|L08611.1|STRMLTODX [153722] gi|153702|gb|J01796.1|STRMALMXP [153702] gij153701|gb|J01795.1|STRMALMX [153701] gi|153693|gb|M13812.1|STRLYTPN [153693] gi|153691|gb|M17717.1|STRLYS [153691] gi|153667|gb|M25525.1|STRKAG73 [153667] gi|398102|gb|L20564.1|STREXP9B [398102] gi|398100|gb|L20563.1|STREXP9A [398100] gi|398098|gb|L20562.1|STREXP8A [398098] gi|398096|gb|L20561.1|STREXP7A [398096] gi|398094|gb|L20560.1|STREXP6A [398094] gi|398092|gb|L20559.1|STREXP5A [398092] gi|398090|gb|L20558.1|STREXP4A [398090] gi|153626|gb|J04234.1|STREXOA [153626] gi|153612|gb|M11226.1|STRDPNM [153612] gi|153603|gb|M25521.1|STRDN87669 [153603] gi|153601|gb|M25526.1|STRDN87577 [153601] gi|153599|gb|M25522.1|STRDN179 [153599] gi|153594|gb|M37688.1|STRDACA [153594] gi|153582|gb|L07752.1|STRATTB [153582] gi|466514|gb|L31413.1|STR1RRA [466514] gi|153551|gb|M25520.1|STR8249 [153551] gi|153549|gb|M25524.1|STR5313972 [153549] gi|153547|gb|M25517.1|STR29044 [153547] gi|153545|gb|M25523.1|STR181071 [153545] gi|153541|gb|M25518.1|STR121 [153541] gi|153539|gb|M25516.1|STR110K70 [153539] gi|506632|gb|U04047.1|SPU04047 [506632] gi|393267|gb|L19055.1|STRPAPA [393267] gi|442066|gb|S62272.1|S62272 [442066] gi|295191|gb|L15190.1|STRPURISYN [295191]

### **CLAIMS**

### What is claimed is:

1. A method for identifying a bacteriophage coding region encoding a product active on an essential bacterial target, comprising identifying a nucleic acid sequence encoding a gene product which provides a bacteria-inhibiting function when said bacteriophage infects a host bacterium,

wherein said bacteriophage is uncharacterized and said host bacterium is a pathogenic bacterium.

- 2. The method of claim 1, further comprising expressing a recombinant bacteriophage ORF in cells of a bacterial strain, wherein inhibition of said cells following expression of said ORF is indicative that said product is active on an essential bacterial target.
- 3. The method of claim 2, wherein inhibition of said bacterium following expression of said ORF is determined by comparison with the growth or viability of said bacterium following expression of an inactivated mutant form of said ORF or in the absence of expression of said ORF, and wherein inhibition of said bacterium following expression of said ORF is indicative that said product is active on an essential bacterial target.
  - 4. The method of claim 2, wherein expression of said ORF is inducible.
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- 5. The method of claim 1, further comprising sequencing at least a portion of a bacteriophage genome.
- 6. The method of claim 1, wherein at least a portion of the nucleotide sequence of a bacteriophage genome is known, said method further comprising identifying at least one ORF in said portion by computer analysis of said sequence.
- 7. The method of claim 6, further comprising analyzing the sequence of said at least one ORF or of a polypeptide encoded by said ORF to identify
   35 homologous genes or gene products of known biochemical function, thereby-indicating the biochemical function of said polypeptide.

- 8. The method of claim 7, wherein said homologous gene or gene product is a bacterial gene important for cell viability.
- 9. The method of claim 7, wherein said homologous gene or gene product 5 is a gene or gene product known to have a bacteria-inhibiting function.
  - 10. The method of claim 6, further comprising analyzing the sequence of said at least one ORF or of a polypeptide encoded by said ORF to identify structural motifs in said polypeptide, thereby indicating the cellular function of said polypeptide.

11. The method of claim 1, wherein a host bacterium for said bacteriophage is selected from the species group consisting of bacteria listed in Table 1.

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- 15 12. The method of claim 1, wherein said bacteriophage is selected from the group consisting of uncharacterized bacteriophage listed in Table 1.
  - 13. The method of claim 2, wherein a plurality of bacteriophage ORFs are expressed in at least one bacterium.
    - 14. The method of claim 13, wherein each of said plurality of bacteriophage ORFs is expressed in a different bacterium.
- 15. The method of claim 14, wherein said plurality of bacteriophage ORFs comprises at least 10% of the ORFs in the genome of said bacteriophage.
  - 16. The method of claim 1, wherein said pathogenic bacterium is an animal pathogen.
- The method of claim 16, wherein said pathogenic bacterium is a human pathogen.
  - 18. The method of claim 1, wherein said pathogenic bacterium is a plant pathogen.
  - 19. The method of claim 1, further comprising confirming the inhibitor function of said ORF.

WO 00/32825 PCT/IB99/02040

- 20. The method of claim 19, wherein said confirming comprises expressing a loss-of-function mutant form of said ORF in said host bacterium.
- 5 21. The method of claim 1, wherein said identifying a nucleic acid sequence encoding a gene product active on an essential bacterial target comprises identifying a nucleic acid sequence encoding a homolog of a bacteriophage polypeptide known to be active on an essential bacterial target.
- 10 22. The method of claim 1, wherein said identifying a bacteriophage coding region comprises identifying a first coding region from a bacteriophage having a non-pathogenic host bacterial strain related to said pathogenic bacterium, said first coding region encoding a product active on an essential bacterial target; and identifying a homolog of said first coding region, wherein said homolog is a probable said bacteriophage coding region encoding a product active on an essential bacterial target.
  - 23. The method of claim 2, wherein a plurality of bacteriophage ORFs from a plurality of different bacteriophage are expressed in at least one bacterium.
  - 24. The method of claim 23, wherein each of said plurality of bacteriophage ORFs are expressed in different bacteria.

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- 25. A method for identifying a target for antibacterial agents, comprising determining the bacterial target of an uncharacterized bacteriophage inhibitor protein.
  - 26. The method of claim 25, wherein said determining comprises identifying at least one bacterial protein which binds to said bacteriophage inhibitor protein or a fragment thereof.
    - 27. The method of claim 26, wherein said binding is determined using affinity chromatography on a solid matrix.
- The method of claim 25, wherein said determining comprises identifying at least one protein:protein interaction using a genetic screen.

- 29. The method of claim 28, wherein said genetic screen is a yeast two-hybrid screen.
- 30. The method of claim 25, wherein said determining comprises a coimmunoprecipitation assay or a protein-protein crosslinking assay.
  - 31. The method of claim 25, wherein said determining comprises identifying a mutated bacterial coding sequence which protects a bacterium from said bacteriophage inhibitor.

- 32. The method of claim 25, wherein said determining comprises identifying a bacterial coding sequence which protects a bacterium against said bacteriophage inhibitor when expressed at high levels in said bacterium.
- 15 33. The method of claim 25, wherein said determining further comprises identifying a bacterial nucleic acid sequence encoding a polypeptide target of said bacteriophage inhibitor protein.
- 34. The method of claim 33, wherein said nucleic acid sequence is
  identified by determining at least a portion of the amino acid sequence of a bacterial protein target, and identifying a bacterial nucleic acid sequence which encodes said protein target.
- 35. The method of claim 25, wherein said bacterial target is naturally
   produced by a bacterial species selected from the group consisting of species of the genera listed in Table 1.
  - 36. The method of claim 25, wherein said bacterial target is naturally produced by a bacterial strain selected from the group consisting of species listed in Table 1.
  - 37. The method of claim 25, wherein said inhibitor protein is naturally produced by a bacteriophage selected from the group consisting of uncharacterized bacteriophage listed in Table 1.

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38. The method of claim 25, further comprising identifying a bacteriophage ORF which encodes a product having a bacteria-inhibiting function.

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- 39. The method of claim 38, wherein said identifying a phage ORF comprises expressing at least one bacteriophage ORF in a bacterium, wherein inhibition of said bacterium following said expression is indicative that said ORF encodes a bacteria-inhibiting function.
- 40. The method of claim 39, wherein a plurality of bacteriophage ORFs are expressed in at least one bacterium.
- 10 41. The method of claim 40, wherein each of said plurality of bacteriophage ORFs is expressed in a different bacterium.
  - 42. The method of claim 41, wherein said plurality of bacteriophage ORFs comprises at least 10% of the ORFs in the genome of said bacteriophage.
  - 43. The method of claim 25, wherein said determining the bacterial target of a bacteriophage inhibitor protein is performed for a plurality of different bacteriophage of the same host bacterium.
- 20 44. The method of claim 25, wherein said bacterial target originates from an animal pathogen.
  - 45. The method of claim 44, wherein said bacterial target is a gene homologous to a gene from an animal pathogen.
    - 46. The method of claim 44, wherein said pathogen is a human pathogen.
  - 47. The method of claim 25, wherein said bacterial target originates from a plant pathogen.
  - 48. The method of claim 25, wherein said bacterial target is a gene homologous to a gene from a plant pathogen.
- 49. The method of claim 25, further comprising determining the cellular or biochemical function or both of said inhibitor protein.

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- 50. The method of claim 25, wherein said identifying the bacterial target comprises identifying a phage-specific site of action.
- 5 51. An isolated, purified, or enriched nucleic acid sequence at least 15 nucleotides in length, wherein said sequence corresponds to at least a portion of a bacteriophage sequence, and wherein said bacteriophage is selected from the group consisting of Staphylococcus aureus bacteriophage 77, 3A, 96, and 44AHJD, Enterococcus baceriophage 182, and Streptococcus pheumoniae bacteriophage Dp-1.

52. The nucleic acid sequence of claim 51, wherein said sequence comprises at least 50 nucleotides.

- 53. The nucleic acid sequence of claim 51, wherein said nucleic acid sequence corresponds to at least a portion of a nucleic acid sequence which encodes a product which provides a bacteria-inhibiting function.
  - 54. The nucleic acid sequence of claim 53, wherein said nucleic acid sequence encodes a polypeptide which provides a bacteria-inhibiting function.
  - 55. The nucleic acid sequence of claim 54, wherein said nucleic acid sequence is transcriptionally linked with regulatory sequences enabling induction of expression of said sequence.
  - 56. An isolated, purified, or enriched polypeptide comprising at least a portion of a protein providing a bacteria-inhibiting function, wherein said polypeptide is normally encoded by a bacteriophage selected from the group consisting of Staphylococcus aureus bacteriophage 77, 3A, 96, and 44AHJD, Enterococcus baceriophage 182, and Streptococcus pheumoniae bacteriophage Dp-1.
  - 57. The polypeptide of claim 56, wherein said polypeptide provides said bacteria-inhibiting function.
- 58. The polypeptide of claim 56, wherein said polypeptide comprises a portion at least 10 amino acid residues in length of a said polypeptide normally encoded by said bacteriophage.

- 59. A recombinant vector comprising a bacteriophage ORF corresponding to an ORF from a bacteriophage having a pathogenic bacterial host, wherein said
   5 bacterial host is selected from the group consisting of uncharacterized bacteria of Table 1.
  - 60. The vector of claim 59, wherein said vector is an expression vector.
- 10 61. The vector of claim 59, wherein said bacteriophage is selected from the group consisting of uncharacterized bacteriophage of Table 1.
  - 62. The vector of claim 61, wherein said bacteriophage is selected from the group consisting of *Staphylococcus aureus* bacteriophage 77, 3A, 96, and 44AHJD, *Enterococcus* baceriophage 182, and *Streptococcus pheumoniae* bacteriophage Dp-1.
    - 63. The vector of claim 60, wherein expression of said ORF is inducible.
- 20 64. A recombinant cell comprising a vector, wherein said vector comprises an ORF from a bacteriophage having a pathogenic bacterial host, wherein said bacterial host is selected from the group consisting of bacterial species of Table 1.
- 65. The recombinant cell of claim 64, wherein said bacteriophage is selected from the group consisting of uncharacterized phage of Table 1.
  - 66. The cell of claim 65, wherein said bacteriophage is selected from the group consisting of *Staphylococcus aureus* bacteriophage 77, 3A, 96, and 44AHJD, *Enterococcus* baceriophage 182, and *Streptococcus pheumoniae* bacteriophage Dp-1.
  - 67. The cell of claim 64, wherein said vector is an expresssion vector and expression of said ORF is inducible.
- 35 68. A method for identifying an antibacterial agent, comprising identifying an active portion of a product of a bacteria-inhibiting ORF of a bacteriophage.

69. The method of claim 68, further comprising constructing a synthetic peptidomimetic molecule, wherein the structure of said molecule corresponds to the structure of said active portion.

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70. A method for identifying a compound active on a target of a bacteriophage inhibitor protein, comprising the step of

contacting a bacterial target protein with a test compound; and
determining whether said compound binds to or reduces the level of
activity of said target protein,

wherein binding of said compound with said target protein or a reduction of the level of activity of said protein is indicative that said compound is active on said target and wherein said target is uncharacterized.

- 71. The method of claim 70, wherein said contacting is carried out in vitro.
- 72. The method of claim 70, wherein said contacting is carried out *in vivo* in a cell.
- The method of claim 70, wherein said compound is a small molecule.
  - 74. The method of claim 70, wherein said compound is a peptidomimetic compound.
- The method of claim 70, wherein said compound is a fragment of a bacteriophage inhibitor protein.
  - 76. The method of claim 70, further comprising determining the site of action of said compound on said target protein.

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77. The method of claim 70, wherein said contacting is performed for a plurality of said target proteins.

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78. A method of screening for potential antibacterial agents, comprising the step of determining whether any of a plurality of compounds is active on a target of a bacteriophage inhibitor protein,

wherein said target is naturally produced by a pathogenic bacterium.

- 79. The method of claim 78, wherein said plurality of compounds are small molecules.
- 80. The method of claim 78, wherein said determining is performed for a plurality of said targets.
- 10 81. A method for inhibiting a bacterium, comprising the step of; contacting said bacterium with a compound active on a target of a bacteriophage inhibitor protein, wherein said target or the target site is uncharacterized.
- 15 82. The method of claim 81, wherein said compound is said protein or an active fragment thereof.
  - 83. The method of claim 81, wherein said compound is a structural mimetic of said protein.
    - 84. The method of claim 81, wherein said compound is a small molecule.
    - 85. The method of claim 81, wherein said contacting is performed in vitro.
- 25 86. The method of claim 81, wherein said contacting is performed *in vivo* in an animal.
  - 87. The method of claim 86, wherein said animal is a human.
- 30 88. The method of claim 81, wherein said contacting is carried out *in vivo* in a plant.
  - 89. The method of claim 81, wherein said bacterium is selected from the group of bacteria listed in Table 1.

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- 90. A method for treating a bacterial infection in an animal suffering from an infection, comprising administering to said animal a therapeutically effective amount of compound active on a target of a bacteriophage inhibitor protein in a bacterium involved in said infection,
- 5 wherein said target is an uncharacterized target or the compound is active at an uncharacterized target site.
  - 91. The method of claim 90, wherein said compound is a small molecule.
- 10 92. The method of claim 90, wherein said compound is a peptidomimetic compound.
  - 93. The method of claim 90, wherein said compound is a fragment of a bacteriophage inhibitor protein.
    - 94. The method of claim 90, wherein said animal is a mammal.
    - 95. The method of claim 94, wherein said mammal is a human.
- 20 96. The method of claim 90, wherein said bacterium is selected from the group listed in Table 1.
  - 97. The method of claim 90, wherein said bacteriophage inhibitor protein is from a bacteriophage selected from the group of bacteriophage listed in Table 1.
  - 98. A method for propylactically treating an animal at risk of an infection, comprising administering to said animal a prophylactically effective amount of a compound active on a target of a bacteriophage inhibitor protein,
- wherein said target is an uncharacterized target or the site of action of said compound is an uncharacterized target site.
  - 99. The method of claim 98, wherein said compound is a small molecule.
- 35 100. The method of claim 98, wherein said compound is a peptidomimetic compound.

- 101. The method of claim 98, wherein said compound is a fragment of a bacteriophage inhibitor protein.
  - 102. The method of claim 98, wherein said animal is a mammal.

- 103. The method of claim 102, wherein said mammal is a human.
- 104. An antibacterial agent active on a target of a bacteriophage inhibitor
  10 protein, wherein said target is an uncharacterized target or said agent is active at a
  phage-specific site on said target.
  - 105. The agent of claim 104, wherein said agent is a pepetidomimetic of a bacteriophage inhibitor polypeptide.

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- 106. The agent of claim 104, wherein said agent is a small molecule.
- 107. The agent of claim 104, wherein said agent is a fragment of a bacteriophage inhibitor polypeptide.

- 108. The agent of claim 104, wherein said agent is active at a phage-specific site on said target.
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- 109. A method of making an antibacterial agent, comprising the steps of:
  - a) identifying a target of a bacteriophage inhibitor polypeptide;
- b) screening a plurality of test compounds to identify a compound active on said target; and
- c) synthesizing said compound in an amount sufficient to provide a
   therapeutic effect when administered to an organism infected by a bacterium naturally producing said target.
  - 110. The method of claim 109, wherein said compound is a small molecule.
- 35 111. The method of claim 109, wherein said compound is a peptidemimetic compound.

- 112. The method of claim 109, wherein said compound is a fragment or derivative of a bacteriophage inhibitor protein.
- 5 113. A computer readable device having recorded therein a nucleotide sequence of a portion of at least one bacteriophage genome of Staphylococcus aureus bacteriophage 77, bacteriophage 3A, or bacteriophage 96, a nucleotide sequence at least 95% identical to a said nucleotide sequence, a ribonucleic acid equivalent, a degenerate equivalent, a homologous sequence, or at least one amino acid sequence encoded by said nucleotide sequence; and

a nucleotide sequence or amino acid sequence analysis program, wherein said program can perform at least one sequence analysis on said nucleotide or amino acid sequence.

- 15 114. The device of claim 113, wherein said at least a portion of at least one bacteriophage genome comprises at least one ORF.
  - 115. The device of claim 113, wherein said device comprises a medium selected from the group consisting of floppy disk, computer hard drive, optical disk, computer random access memory, and magnetic tape wherein said nucleotide or amino acid sequence or said program or both are recorded on said medium.
  - 116. The device of claim 113, wherein said portion of at least one bacteriophage genomic nucleotide sequence comprises at least 50% of at least one bacteriophage genomic sequence.
  - 117. The device of claim 113, wherein said at least one bacteriophage nucleotide genomic sequence comprises portions of a plurality of bacteriophage nucleotide genomic sequences.

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- 118. A computer-based system for identifying biologically important portions of a bacteriophage genome, comprising:
- a) a data storage medium having recorded thereon a nucleotide sequence
  corresponding to a portion of at least one bacteriophage genome, wherein said
  bacteriophage genome is uncharacterized;

- b) a set of instructions allowing searching of said sequence to analyze said sequence; and
  - c) an output device.

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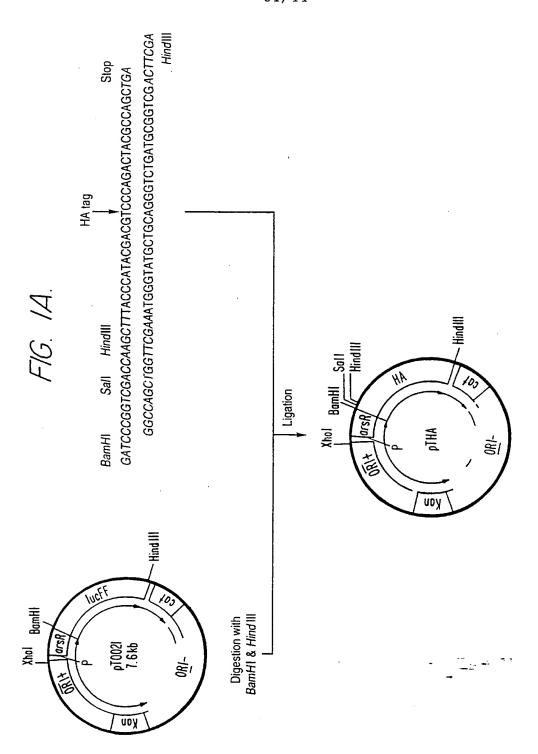
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- The system of claim 118, wherein said output device comprises comprises a device selected from the group consisting of a printer, a video display, and a recording medium.
- 120. The system of claim 118, wherein said bacteriophage genome is of a bacteriophage selected from the group consisting of uncharacterized bacteriophage listed in Table 1.
  - 121. The system of claim 118, wherein said uncharacterized bacteriophage is selected from the group consisting of bacteriophage 77, 3A, and 96.

122. A method for identifying or characterizing a bacteriophage ORF, comprising the steps of:

- a) providing a computer-based system for analyzing nucleic acid or amino acid sequence data, wherein said system comprises a data storage medium having recorded thereon at least one nucleotide or amino acid sequence corresponding to a portion of at least one uncharacterized bacteriophage genome, a set of instructions allowing searching of said sequence to analyze said sequence; and an output device;
  - b) analyzing at least a portion of at least one said sequence; and
  - c) outputting results of said analyzing to said output device.
- 123. The method of claim 122, wherein said analysis identifies sequence similarity or homology with sequences selected from the group consisting of bacterial ORFs encoding products with related biological function; ORFs encoding known inhibitors or bacteria, essential bacterial ORFs.
- 124. The method of claim 122, wherein said analysis comprises identifying a probable biological function based on identification of structural elements or sequence homology or similarity.
- 125. The method of claim 122, wherein said bacteriophage is selected from the group consisting of uncharacterized bacteriophage listed in Table 1.

126. The method of claim 125, wherein said uncharacterized bacteriophage is selected from bacteriophage 77, 3A, and 96.



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TGA GAAAAGGAGGCGGATCCATG

BamHI

arsR

CTCGAG-

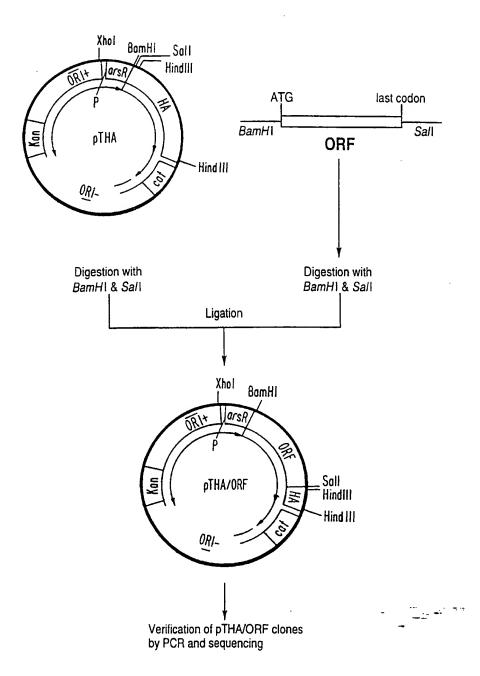
Xhol

LucFF

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Hind III

FIG. 2.

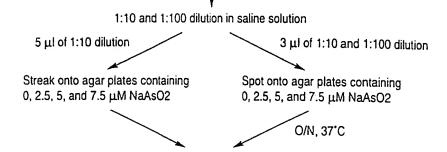


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FIG. 3.

#### (A) Functional assay on semi-solid support media

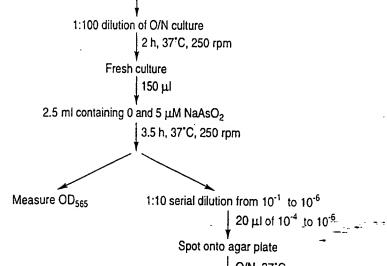
Frozen stock of phage 77 pTHA/ORF S. aureus RN4220 transformants



Compare bacterial growth on plates with and without NaAsO2

#### (B) Functional assay in liquid medium

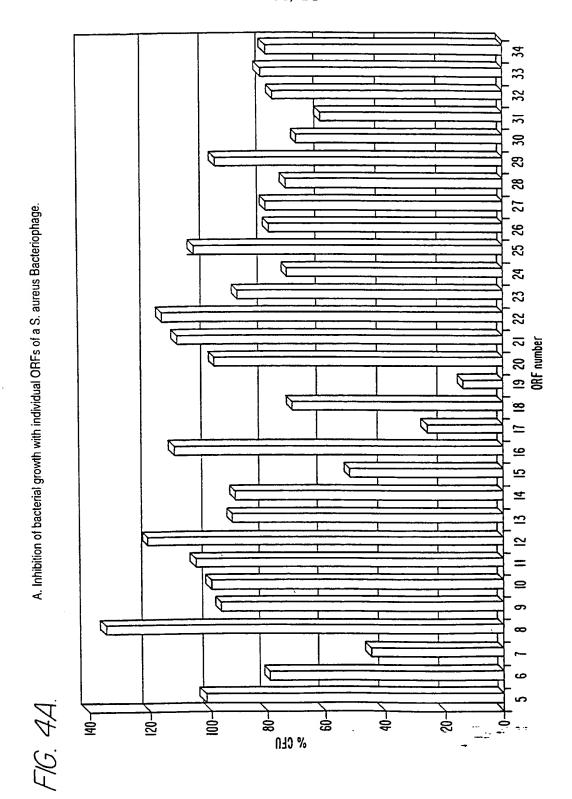
O/N culture inoculated from frozen stock of phage 77 pTHA/ORF *S. aureus* RN4220 transformants



Count colonies

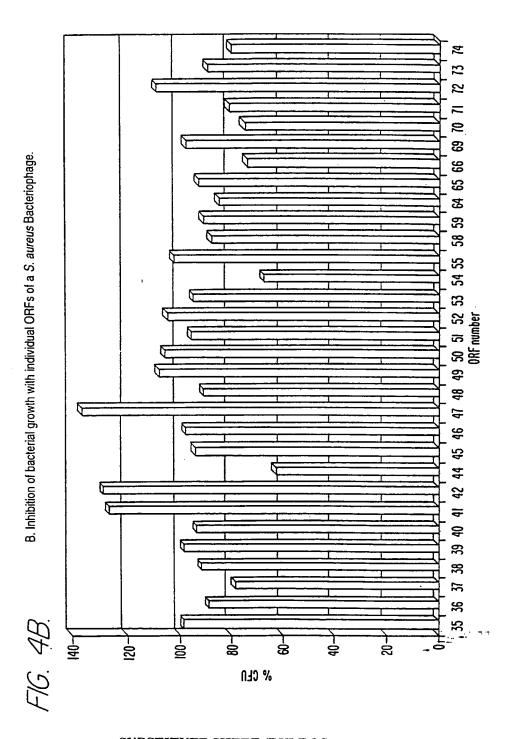
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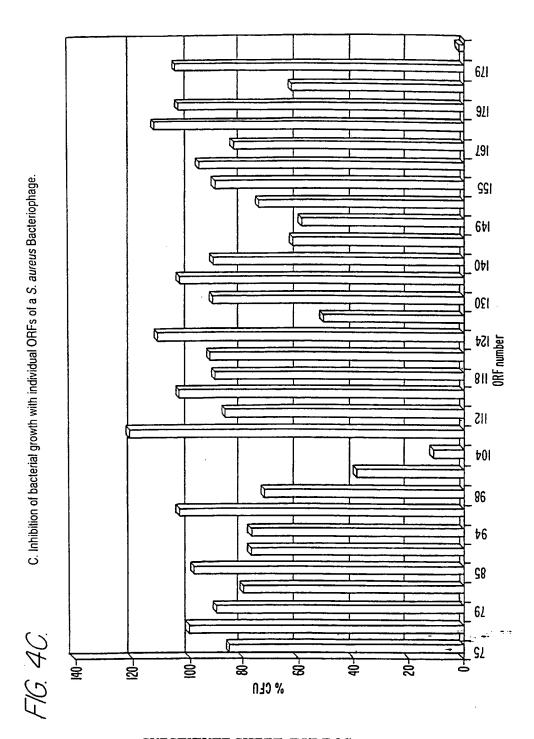
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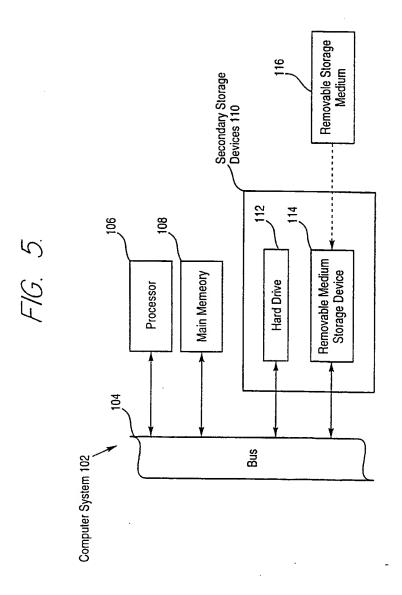


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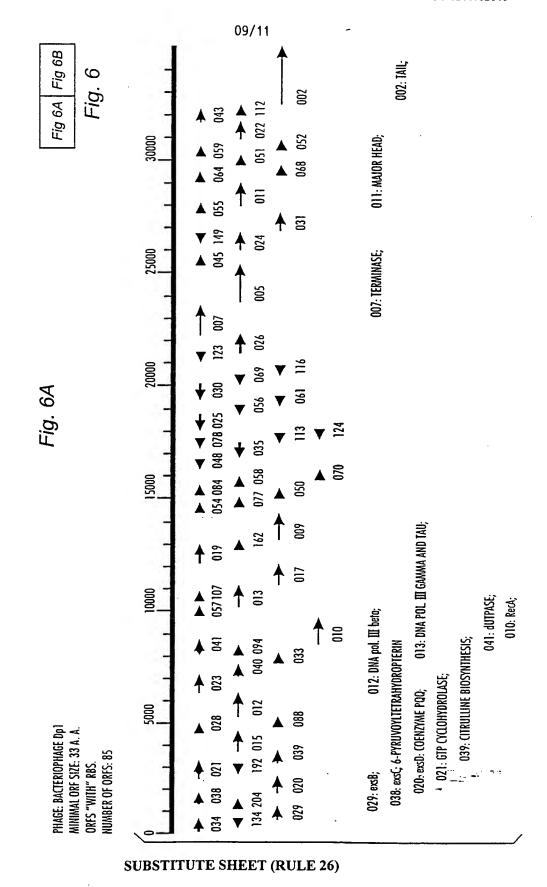
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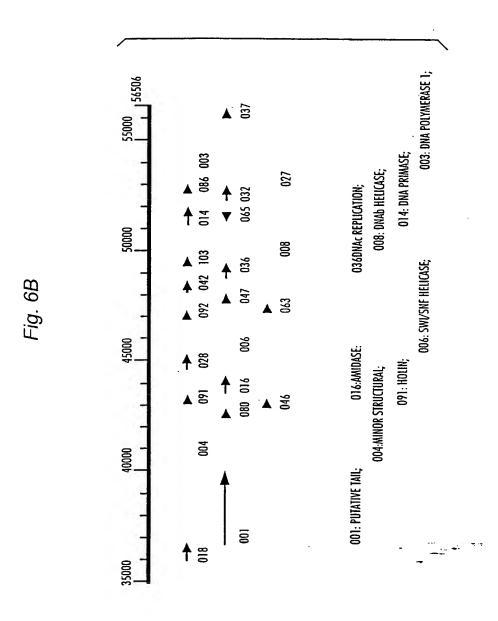


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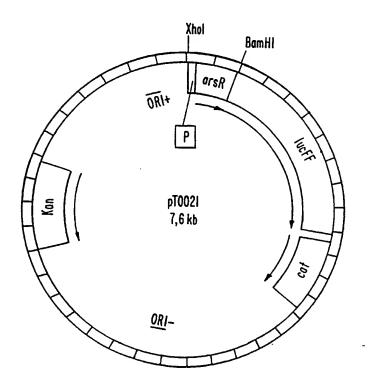
# FIG. 7.

#### Abbreviations:

kan: gene encoding kanamycin resistance cat: gene encoding chloramphenicol resistance ori + and -: origin of replication in gram-positive and gram-negative bacteria, respectively arsR: gene encoding regulatory protein of the ars promoter P: ars promoter lucFF: gene encoding luciferase protein. This portion will be removed and replaced by individual *S. aureus* phage genes.

#### Referance:

Tauriainen et al., Appl. Environ. Microbio. 1997. 63: 4456-4461



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# (19) World Intellectual Property Organization International Bureau



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2 December 1999 (02.12.1999)

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09/454,252

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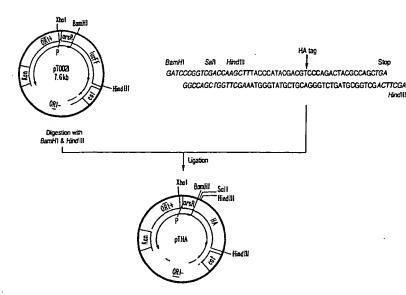
Jerry [CA/CA]; 8 Lakeview, Baie D'Urfe, Quebec H9X 3B1 (CA). GROS, Phillippe [CA/CA]; 107 Montrose, St. Lambert, Quebec J4R 1X4 (CA). DUBOW, Michael [CA/CA]; 4901 Coolbrook Avenue, Montreal, Quebec H3X 2K8 (CA).

- (74) Agents: MORROW, Joy, D. et al.; Smart & Biggar, 900 - 55 Metcalfe Street, P.O. Box 2999, Station D, Ottawa, Ontario K1P 5Y6 (CA).
- (81) Designated States (national): AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW.
- (84) Designated States (regional): ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).

[Continued on next page]

#### (54) Title: DEVELOPMENT OF ANTI-MICROBIAL AGENTS BASED ON BACTERIOPHAGE GENOMICS

US



00/32825 A3

(57) Abstract: A method for identifying suitable targets for antibacterial agents based on identifying targets of bacteriophage-encoded proteins is described. Also described are compositions useful in the identification methods and in inhibiting bacterial growth, and methods for preparing and using such compositions.



#### Published:

- With international search report.
- (88) Date of publication of the international search report: 18 January 2001

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

#### INTERNATIONAL SEARCH REPORT

Intern. .nal Application No PCT/IB 99/02040

PCT/IB 99/02040 A. CLASSIFICATION OF SUBJECT MATTER IPC 7 C12Q1/70 C12Q1/68 C12N15/10 C12N15/34 C12N1/21 C07K14/01 C12Q1/18 According to international Patent Classification (IPC) or to both national classification and IPC B. FIELDS SEARCHED Minimum documentation searched (classification system followed by classification symbols) C120 C12N C07K Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched Electronic data base consulted during the international search (name of data base and, where practical, search terms used) C. DOCUMENTS CONSIDERED TO BE RELEVANT Relevant to claim No. Citation of document, with indication, where appropriate, of the relevant passages Category ° EP 0 072 925 A (RUTGERS RES & EDUCATION X 1,2, FOUND) 2 March 1983 (1983-03-02) 11-18 the whole document 3-5,19,Υ 20,22-24 SHEEHAN, M.M. ET AL.: "The lytic enzyme 1,2,11, X of the pneumococcal phage Dp-1: a chimeric 12,16,17 lysin of intergeneric origin." MOLECULAR MICROBIOLOGY, vol. 25, no. 4, 1997, pages 717-25, XP000922620 Y the whole document 3-5,19,20,22-24 -/--Further documents are listed in the continuation of box C. Patent family members are listed in annex. X Special categories of cited documents : "T" later document published after the international filing date or priority date and not in conflict with the application "A" document defining the general state of the art which is not considered to be of particular relevance cited to understand the principle or theory underlying the "E" earlier document but published on or after the international "X" document of particular relevance; the claimed invention filing date cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the "O" document referring to an oral disclosure, use, exhibition or document is combined with one or more other such docu ments, such combination being obvious to a person skilled other means in the art. \*P\* document published prior to the international filing date but later than the priority date claimed \*&\* document member of the same patent family Date of mailing of the international search report Date of the actual completion of the international search 0 5 10.00

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European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Authorized officer

Smalt, R

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CARL R (US); US GOVERNMENT (US); ADHYA S) 12 October 1995 (1995-10-12)  the whole document  KANEKO J ET AL: "Complete nucleotide sequence and molecular characterization of the temperate staphylococcal bacteriophage phiPVL carrying Panton-Valentine leukocidin genes" GENE,NL,ELSEVIER BIOMEDICAL PRESS. AMSTERDAM, vol. 215, no. 1, pages 57-67, XP004149229 ISSN: 0378-1119 cited in the application the whole document  WO 89 00199 A (UNIV LOUISIANA STATE) 12 January 1989 (1989-01-12)  the whole document  EP 0 748 871 A (NESTLE SA) 18 December 1996 (1996-12-18)	evant to claim No.						
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18 December 1996 (1996-12-18)							

International application No. PCT/IB 99/02040

#### INTERNATIONAL SEARCH REPORT

Box I	Observations where certain claims were found unsearchable (Continuation of Item 1 of first sheet)
This Inte	emational Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:
1. 🔲	Claims Nos.: because they relate to subject matter not required to be searched by this Authority, namely:
2.	Claims Nos.: because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
3.	Claims Nos.: because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).
Box II	Observations where unity of invention is lacking (Continuation of item 2 of first sheet)
This Int	ernational Searching Authority found multiple inventions in this international application, as follows:
1.	As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2.	As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3.	As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. X.	No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:  invention 1: claims 2-4, 13-15, 23, 24 compl. and 1, 5, 11, 12, 16-20, 22, 68, 69 partially
Remar	the additional search fees were accompanied by the applicant's protest.  No protest accompanied the payment of additional search fees.

#### FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

Invention 1: claims 2-4,13-15,23,24 completely, and 1,5,11,12,16-20,22,68,69 partially

Method for identifying bacteriophage-encoded inhibitors of pathogenic bacterial targets by expression of the viral ORF in a host cell.

Invention 2: claims 6-10,21,
 118-126 completely and 1,5,11,12,16-20,22,68,
 69 partially

Method for identifying bacteriophage-encoded inhibitors of pathogenic bacterial targets by computer-based methods, and computer system for use therein.

Invention 3: claims 25-50

Method for identifying the pathogenic bacterial target of a bacteriophage encoded product.

Invention 4: claims 51-67,81-103,113-117, all partially, and as far as applicable

Isolated polynucleotides of at least 15 nucleotides in length corresponding to at least a portion of sequence ID.1, peptides comprising a portion of at least 10 amino acids normally encoded by seq.ID.1, hosts, recombinant production of the protein, computer-readable devices containing sequence data of seq.ID.1, and a method for inhibiting a pathogenic bacterium using the protein encoded by seq.ID.1, all as far as applicable.

Inventions 5-2639 : claims 51-67,81-103,113-117, all partially, and as far as applicable

Idem as invention 4, but limited to the respective seq.ID's 2-2636, whereby invention 5 relates to seq.ID.2, invention 6 relates to seq.ID.3, ....., and invention 2639 relates to seq.ID.2636.

For the sake of conciseness, the general subject matter is explicitly defined, the specific subject matters of each invention are defined by analogy thereto.

Invention 2640: claims 70-80,109-112

#### FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

Method for identifying a compound active on the pathogenic bacterial target of a bacteriophage-encoded inhibitor and method for producing said compound.

Invention 2641: claims 103-108

Antibacterial agents active on the target of phage-encoded inhibitors in pathogenic bacteria.

page 2 of 2

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Information on patent family members

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